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OM protein - protein search, using sw model

Run on: September 24, 2004, 14:02:27 : Search time 51.308 seconds
(without alignments)
659.375 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 687

Sequence: 1 GSGKDPVQPTKICVGRD.....VPWEKKIYTVVNHMECEP 127

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_29Jan04:*

- 1: Geneseq1980s:*
- 2: Geneseq1990s:*
- 3: Geneseq2000s:*
- 4: Geneseq2001s:*
- 5: Geneseq2002s:*
- 6: Geneseq2003s:*
- 7: Geneseq2003bs:*
- 8: Geneseq2004s:*

Prod. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB ID	Description
1	618	90.0	123	3	AA95426 Human hig
2	618	90.0	304	6	ABP70801 Human ext
3	618	90.0	322	6	ABP70799 Human ext
4	618	90.0	329	6	ABU92044 Human pro
5	618	90.0	358	6	ABP70800 Human ext
6	618	90.0	390	6	ABU99149 Novel hum
7	618	90.0	398	6	ABU99143 Novel hum
8	618	90.0	427	8	ABP76864 Human pro
9	618	90.0	615	6	ABU99144 Novel hum
10	618	90.0	626	5	ABP78707 Human hig
11	618	90.0	644	4	ABG21101 Novel hum
12	618	90.0	644	5	ABP78710 Human hig
13	618	90.0	644	6	ABU99150 Novel hum
14	618	90.0	644	6	ABU99145 Novel hum
15	586	85.3	122	3	ABP7447 Human kin
16	585	85.2	435	4	ABG21105 Novel hum
17	556.5	81.0	117	2	AA933350 Domaine 3
18	440	64.0	436	1	AA940257 Bradykini
19	413	60.1	434	1	AA940633 Bradykini
20	411	59.8	357	6	ABR41202 Human DIT
21	388	56.5	235	5	ABG60077 Human DIT
22	370	54.7	248	4	ABG21102 Novel hum
23	316	46.0	369	4	ABG21099 Novel hum
24	190	27.7	305	4	ABG21100 Novel hum
25	171.5	25.0	167	2	AA998907 Mouse IMC

RESULT 1
AA95426
ID AA95426 standard; peptide; 123 AA.

XX AA95426;

DT 25-SEP-2000 (first entry)

XX Human high mol.wt. kininogen domain 3.

XX Anti-angiogenic; angiogenesis; inhibitor; kininogen; homologue;
XX endothelial cell proliferation; apoptosis; cancer; ocular disorder;
XX rheumatoid arthritis; cytostatic; antiarthritic; antirheumatic; therapy;
XX human; D3 peptide.

XX Homo sapiens.

XX WO200035407-A2.

XX 22-JUN-2000.

XX 02-DEC-1999; 99WO-US028465.

XX 16-DEC-1998; 98US-0112427P.

XX (UTEM) UNIV TEMPLE.

XX (MCCR/) MCCRAE R K.

XX Mcrae RK;

XX WPI; 2000-442247/38.

XX Composition for inhibiting angiogenesis and endothelial cell proliferation, inducing endothelial cell apoptosis and treating cancer, rheumatoid arthritis, and ocular disorders comprises a kininogen domain 3 analog.

XX Disclosure, Page 4; 44pp; English.

XX The present sequence is that of domain 3 of human high mol.wt. kininogen (HK). The invention provides peptides (see AA95405-24) that are analogues of certain sites in the HK domain 3, specifically Asn275-Lys282, Cys246-Cys249, Leu331-Tyr338 and Tyr299-Ser314. The peptides, in which native Cys residues may be replaced by Ala residues, inhibit endothelial cell proliferation and may also induce endothelial cell apoptosis. Compositions including the peptides are used in claimed methods for inhibiting angiogenesis, inhibiting endothelial cell proliferation, and inducing endothelial cell apoptosis. Cancer.

ALIGNMENTS

26	166	24.2	32	3	AA95418
27	163.5	23.8	126	3	AA95418
28	163.5	23.8	145	2	AA95418
29	163.5	23.8	145	2	AA95418
30	163.5	23.8	145	2	AA95418
31	163.5	23.8	145	2	AA95418
32	163.5	23.8	145	2	AA95418
33	163.5	23.8	145	2	AA95418
34	163.5	23.8	145	2	AA95418
35	163	23.7	167	7	ADA45154
36	163	23.7	167	7	ADA45154
37	161	23.4	178	2	AA95418
38	157	22.9	122	3	AA95418
39	154.5	22.5	167	2	AA95418
40	153	22.3	27	3	AA95425
41	144.5	21.0	121	3	AA95425
42	144.5	21.0	128	3	AA95425
43	143.5	20.9	121	3	AA95425
44	143.5	20.9	128	3	AA95425
45	142.5	20.7	118	3	AA95425

CC rheumatoid arthritis, and ocular disorders characterized by undesired
 CC vascularization of the retina are treated
 XX
 SQ Sequence 123 AA;
 Query Match 90.0%; Score 618; DB 3; Length 123;
 Best Local Similarity 100.0%; Pred. No. 5.5e-63;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDVPPTKICVCGPRDIPNTPSPLEBETLTHITIKLNAENATFFPKIDNVKARVQV 62
 DB 1 GKDVPPTKICVCGPRDIPNTPSPLEBETLTHITIKLNAENATFFPKIDNVKARVQV 60
 QY 63 AGKYFIDFVARETTCSKESNEELTESCTKGLGSLDCNAEVVVPWEKKIYPTV 118
 DB 61 AGKYFIDFVARETTCSKESNEELTESCTKGLGSLDCNAEVVVPWEKKIYPTV 116
 RESULT 2
 ABP70801
 ID ABP70801 standard; protein; 304 AA.
 XX
 AC ABP70801;
 XX
 DT 26-AUG-2003 (first entry)
 XX
 DE Human extracellular messenger, EXMES-28.
 XX
 KW Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
 KW immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
 KW endocrine disorder; cancer.
 XX
 OS Homo sapiens.
 XX
 PN WO2003018612-A2.
 XX
 PD 06-MAR-2003.
 XX
 PF 22-AUG-2002; 2002WO-US027213.
 XX
 PR 24-AUG-2001; 2001US-0314811P.
 PR 14-DEC-2001; 2001US-0340584P.
 PR 18-JAN-2002; 2002US-0350595P.
 PR 11-MAR-2002; 2002US-0363432P.
 PR 15-MAR-2002; 2002US-0364607P.
 PR 05-APR-2002; 2002US-0370761P.
 PR 24-JUN-2002; 2002US-0391378P.
 XX
 PA (INCY-) INCYTE GENOMICS INC.
 XX
 PI Duggan BM, Lee S, Baughn MR, Hafalia AJA, Wallia NK, Elliott VS,
 PI Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang YT, Burford N;
 PI Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebardjian Y,
 PI Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PM;
 PI Rankumar J;
 XX
 DR WPI; 2003-278643/27.
 DR N-PSDB; ACC42388.
 XX
 XX New human extracellular messenger (EXMES) polypeptide, useful for
 PT preparing a composition for treating a disease associated with decreased
 PT expression or overexpression of functional EXMES e.g. autoimmune
 PT disorders or cancer.
 XX
 PS Claim 1; Page 207; 22app; English.
 XX
 CC The present invention relates to novel human extracellular messenger
 CC proteins (EXMES-1 to-28; ABP70774-ABP70801) and their coding sequences
 CC (ACC42361-ACC42388). The proteins are useful for preparing a composition
 CC for diagnosing or treating a disease or condition associated with
 CC decreased expression or overexpression of functional EXMES e.g.
 CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or
 CC cancer

XX SQ Sequence 304 AA;
 Query Match 90.0%; Score 618; DB 6; Length 304;
 Best Local Similarity 100.0%; Pred. No. 1.9e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDVPPTKICVCGPRDIPNTPSPLEBETLTHITIKLNAENATFFPKIDNVKARVQV 62
 DB 130 GKDVPPTKICVCGPRDIPNTPSPLEBETLTHITIKLNAENATFFPKIDNVKARVQV 189
 QY 63 AGKYFIDFVARETTCSKESNEELTESCTKGLGSLDCNAEVVVPWEKKIYPTV 118
 DB 190 AGKYFIDFVARETTCSKESNEELTESCTKGLGSLDCNAEVVVPWEKKIYPTV 245
 RESULT 3
 ABP70799
 ID ABP70799 standard; protein; 322 AA.
 XX
 AC ABP70799;
 XX
 DT 26-AUG-2003 (first entry)
 XX
 DE Human extracellular messenger, EXMES-36.
 XX
 KW Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
 KW immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
 KW endocrine disorder; cancer.
 XX
 OS Homo sapiens.
 XX
 PN WO2003018612-A2.
 XX
 PD 06-MAR-2003.
 XX
 PF 22-AUG-2002; 2002WO-US027213.
 XX
 PR 24-AUG-2001; 2001US-0314811P.
 PR 14-DEC-2001; 2001US-0340584P.
 PR 18-JAN-2002; 2002US-0350595P.
 PR 11-MAR-2002; 2002US-0363432P.
 PR 15-MAR-2002; 2002US-0364607P.
 PR 05-APR-2002; 2002US-0370761P.
 PR 24-JUN-2002; 2002US-0391378P.
 XX
 PA (INCY-) INCYTE GENOMICS INC.
 XX
 PI Duggan BM, Lee S, Baughn MR, Hafalia AJA, Wallia NK, Elliott VS,
 PI Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang YT, Burford N;
 PI Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebardjian Y,
 PI Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PM;
 PI Rankumar J;
 XX
 DR WPI; 2003-278643/27.
 DR N-PSDB; ACC42386.
 XX
 XX New human extracellular messenger (EXMES) polypeptide, useful for
 PT preparing a composition for treating a disease associated with decreased
 PT expression or overexpression of functional EXMES e.g. autoimmune
 PT disorders or cancer.
 XX
 PS Claim 1; Page 205-206; 22app; English.
 XX
 CC The present invention relates to novel human extracellular messenger
 CC proteins (EXMES-1 to-28; ABP70774-ABP70801) and their coding sequences
 CC (ACC42361-ACC42388). The proteins are useful for preparing a composition
 CC for diagnosing or treating a disease or condition associated with
 CC decreased expression or overexpression of functional EXMES e.g.
 CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or
 CC cancer

Query Match 90.0%; Score 618; DB 6; Length 322;
 Best Local Similarity 100.0%; Pred. No. 2.1e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GNDVPPTKICVGCPRDPTNSPELETLTHITIKLNAENNAFFPKIDNVKARQVV 62
 DB 148 GNDVPPTKICVGCPRDPTNSPELETLTHITIKLNAENNAFFPKIDNVKARQVV 207

QY 63 AGKVFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTV 118
 DB 208 AGKVFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTV 263

RESULT 4
 ABUS92044
 ID ABUS92044 standard; protein; 329 AA.

XX AC ABUS92044;
 XX DT 15-JUL-2003 (first entry)
 XX DE Human protein modification and maintenance molecule-24 (PMM-24).
 XX KW Human; protein modification and maintenance molecule; PMM; cancer;
 KW cell proliferation disorder; atherosclerosis; neurological disorder;
 KW epilepsy; Huntington's disease; stroke; immune disorder; allergy;
 KW inflammatory disorder; AIDS; developmental disorder; hypothyroidism;
 KW Cushing's syndrome; gastrointestinal disorder; epithelial disorder;
 KW infection; cytostatic; antiarteriosclerotic; anticonvulsant; nootropic;
 KW neuroprotective; cerebroprotective; anti-HIV; anti-allergic; vulnary;
 KW antiinflammatory; thyromimetic.

XX OS Homo sapiens.
 XX PN WO2003031939-A2.

XX PD 17-APR-2003.

XX PP 11-OCT-2002; 2002WO-US032850.

XX PR 12-OCT-2001; 2001US-0329689P.

XX PR 25-OCT-2001; 2001US-0335703P.

XX PR 09-NOV-2001; 2001US-0348887P.

XX PR 28-NOV-2001; 2001US-0334145P.

XX PR 06-DEC-2001; 2001US-0337451P.

XX PR 14-DEC-2001; 2001US-0340584P.

XX PA (INCY-) INCYTE GENOMICS INC.

XX PI Rankumar J, Gorvad AE, Baughn MR, Emerling BM, Yang J, Lee SY;
 PI Tran UK, Becha SD, Duggan BM, Lee EA, Griffin JA, Li JX;
 PI Sprague KM, Hafalia AJA, Chawla NK, Lehr-Mason PM, Kable AG, Yue H;
 PI Marquis JP, Yao MG, Richardson TW, Tang TY, Jin P, Chien D;
 PI Bhatia U, Burkill JD, Lee S, Blake JJ, Ho A, Zheng W;

XX DR WPI; 2003-430274/40.

XX DR N-PSDB; ACA92439.

XX PT New human protein modification and maintenance molecules (PMM), useful
 PT for diagnosing, treating and preventing diseases or conditions associated
 PT with the aberrant PMM expression e.g. cancer, atherosclerosis, or
 PT infections.

XX PS Claim 1; Page 264-265; 311pp; English.

XX CC The present invention relates to the isolation of human protein
 CC modification and maintenance molecules (PMM), and the polynucleotide
 CC sequences encoding them. A total of 40 PMM polypeptides (designated PMM
 CC 1 to PMM-40) are disclosed. The sequences of the invention are useful
 CC for diagnosing a condition or disease associated with the expression of
 CC PMM in a subject, preparing a polyclonal or monoclonal antibody, and
 CC generating an expression profile of a sample containing the

CC polynucleotides. The diseases or conditions associated with decreased
 CC expression or overexpression of PMM are cell proliferation disorders
 CC (e.g. cancer, atherosclerosis), neurological disorders (e.g. epilepsy,
 CC Huntington's disease, stroke), immune/inflammatory disorders, (e.g. AIDS,
 CC allergies), developmental disorders (e.g. hypothyroidism, Cushing's
 CC syndrome), gastrointestinal or epithelial disorders, and infections. The
 CC PMM polypeptides or their fragments are useful in screening compounds
 CC for effectiveness as agonists or antagonists of the polypeptides, or in
 CC altering the expression of the target polynucleotide and compounds that
 CC specifically bind to, or modulate the activity of the polypeptide.
 CC ABUS92021-ABUS92060 represent the human PMM polypeptides of the invention
 XX
 XX Sequence 329 AA;

Query Match 90.0%; Score 618; DB 6; Length 329;

Best Local Similarity 100.0%; Pred. No. 2.2e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GNDVPPTKICVGCPRDPTNSPELETLTHITIKLNAENNAFFPKIDNVKARQVV 62
 DB 155 GNDVPPTKICVGCPRDPTNSPELETLTHITIKLNAENNAFFPKIDNVKARQVV 214

QY 63 AGKVFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTV 118

DB 215 AGKVFIDFVARETTCSKESNEELTESCETKLGSLDCNAEVVVPWEKKIYPTV 270

RESULT 5

ABP70800

ID ABP70800 standard; protein; 359 AA.

XX AC ABP70800;

XX DT 26-AUG-2003 (first entry)

XX DE Human extracellular messenger, EXMES-37.

XX KW Human; extracellular messenger; EXMES; cytostatic; antidiabetic;
 KW immunosuppressive; autoimmune disorder; inflammatory disorder; diabetes;
 KW endocrine disorder; Cancer.

XX OS Homo sapiens.

XX PN WO2003018612-A2.

XX PD 06-MAR-2003.

XX PP 22-AUG-2002; 2002WO-US027213.

XX PR 24-AUG-2001; 2001US-0314811P.

XX PR 14-DEC-2001; 2001US-0340584P.

XX PR 18-JAN-2002; 2002US-0350595P.

XX PR 11-MAR-2002; 2002US-0363432P.

XX PR 15-MAR-2002; 2002US-0364607P.

XX PR 05-APR-2002; 2002US-0370761P.

XX PR 24-JUN-2002; 2002US-0391378P.

XX PA (INCY-) INCYTE GENOMICS INC.

XX PI Duggan BM, Lee S, Baughn MR, Hafalia AJA, Wallia NK, Elliott VS;
 PI Lee SY, Khare R, Wilson AD, Jin P, Hawkins PR, Tang YT, Burford N;
 PI Ding L, Yao MG, Becha SD, Tran UK, Chien D, Zebajadian Y;
 PI Richardson TW, Kable AE, Chang H, Swarnakar A, Lehr-Mason PM;
 PI Rankumar J;

XX DR WPI; 2003-278643/27.

XX DR N-PSDB; ACC42387.

XX CC New human extracellular messenger (EXMES) polypeptide, useful for
 CC preparing a composition for treating a disease associated with decreased
 CC expression or overexpression of functional EXMES e.g. autoimmune
 CC disorders or cancer.

PS Claim 1; Page 206; 224pp; English.

XX The present invention relates to novel human extracellular messenger

CC proteins (EXMES-1 to -28; ABP70774-ABP70801) and their coding sequences

CC (ACC42361-ACC42388). The proteins are useful for preparing a composition

CC for diagnosing or treating a disease or condition associated with

CC decreased expression or overexpression of functional EXMES e.g.

CC autoimmune/inflammatory disorders, diabetes, endocrine disorders or

CC cancer

XX Sequence 358 AA;

XX

Query Match 90.0%; Score 618; DB 6; Length 358;

Best Local Similarity 100.0%; Pred. No. 2.4e-62;

Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 GKDFVQPPPTKICVGCPRDPTNSPELEETLTHITKLNAENNAATPFKIDNVKKARVQV 62

DB 184 GKDFVQPPPTKICVGCPRDPTNSPELEETLTHITKLNAENNAATPFKIDNVKKARVQV 243

OY 53 AGKKYFIDFVARETTCKESNEELTSCETKLGQSLDCNAEVYVVPWEKKIYPTV 118

DB 244 AGKKYFIDFVARETTCKESNEELTSCETKLGQSLDCNAEVYVVPWEKKIYPTV 299

RESULT 6

ABU99149

ID ABU99149 standard; protein; 390 AA.

AC ABU99149;

XX

XX 01-AUG-2003 (first entry)

XX Novel human GPCR related protein NOV12g.

XX Human; G-protein coupled receptor related protein; GPCR related protein;

XX NOV; cytotactic; cardiac; antiarteriosclerotic; antidiabetic;

XX immunomodulator; anti-HIV; anorectic; antisthmatic; haemostatic;

XX antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;

XX NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;

XX diabetes; immune disorder; AIDS; obesity; asthma;

XX haematopoietic disorder; Parkinson's disease; Alzheimer's disease;

XX infection; multiple sclerosis; cancer-associated cachexia;

XX wasting disorder; chronic disease; neurogenesis; cell differentiation;

XX cell proliferation; haematopoiesis; wound healing; angiogenesis;

XX chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.

XX Homo sapiens.

XX

XX WO2002299116-A2.

XX

XX 12-DEC-2002.

XX

XX 04-JUN-2002; 2002WO-US017428.

XX

XX 04-JUN-2001; 2001US-0295607P.

XX

XX 04-JUN-2001; 2001US-0295661P.

XX

XX 06-JUN-2001; 2001US-0296404P.

XX

XX 06-JUN-2001; 2001US-0296418P.

XX

XX 14-JUN-2001; 2001US-0298285P.

XX

XX 15-JUN-2001; 2001US-0298554P.

XX

XX 21-JUN-2001; 2001US-0299949P.

XX

XX 26-JUN-2001; 2001US-0300881P.

XX

XX 28-JUN-2001; 2001US-0301509P.

XX

XX 13-AUG-2001; 2001US-0311972P.

XX

XX 29-AUG-2001; 2001US-0315607P.

XX

XX 14-SEP-2001; 2001US-0322291P.

XX

XX 17-SEP-2001; 2001US-0322706P.

XX

XX 14-DEC-2001; 2001US-0341186P.

XX

XX 28-FEB-2002; 2002US-0361189P.

XX

XX 12-MAR-2002; 2002US-0363673P.

XX

XX 12-MAR-2002; 2002US-0363676P.

PR 03-JUN-2002; 2002US-00363676.

XX (CURA-) CURAGEN CORP.

XX Anderson DW, Baumgartner JC, Boldog PL, Casman SJ, Edinger SR;

XX Gargoli EA, Gerlach VL, Gorman L, Guo X, Hjalt T, Kekuda R, Li L;

XX MacDougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M;

XX Pena CEA, Rastelli L, Shinkens RA, Stone DJ, Spytek KA, Vernet CAM;

XX Voss EZ, Zerhusen BD;

XX WPI; 2003-140627/13.

XX N-ESDB; ACD03653.

XX New NOVX polypeptides and nucleic acids, useful for preventing or

XX treating NOVX-associated disorders e.g. cancer, cardiomyopathy,

XX atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or

XX pharmacogenomics.

XX Claim 1; Page 147; 332pp; English.

XX The invention describes an isolated polypeptide (I) comprising any of 27

XX 118-961 residue amino acid sequences, given in the specification, a

XX mature form of them, a sequence that is at least 95 % identical to them,

XX or a sequence having one or more conservative substitutions in them. The

XX polypeptide is useful in manufacturing a medicament for treating a

XX syndrome associated with a human disease selected from a pathology

XX associated with the polypeptide. The NOVX polypeptides, polynucleotides

XX and antibodies are useful in treating or preventing NOVX-associated

XX disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune

XX disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's

XX disease, Alzheimer's disease, infections, multiple sclerosis, cancer-

XX associated cachexia, and other wasting disorders associated with chronic

XX diseases. The nucleic acids and polypeptides may also be used as targets

XX for the identification of small molecules that modulate or inhibit e.g.

XX neurogenesis, cell differentiation, cell proliferation, haematopoiesis,

XX wound healing and angiogenesis, in gene therapy, in generation of

XX antibodies that bind immunospecifically to NOVX substances for use in

XX therapeutic or diagnostic methods. The nucleic acids are further used as

XX hybridisation probes, in chromosome mapping, tissue typing, preventive

XX vaccines, and pharmacogenomics. The polypeptides are also useful as

XX medicines. This is the amino acid sequence of a novel human G-protein

XX coupled receptor related protein NOV

XX Sequence 390 AA;

XX

Query Match 90.0%; Score 618; DB 6; Length 390;

Best Local Similarity 100.0%; Pred. No. 2.7e-62;

Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 GKDFVQPPPTKICVGCPRDPTNSPELEETLTHITKLNAENNAATPFKIDNVKKARVQV 62

DB 216 GKDFVQPPPTKICVGCPRDPTNSPELEETLTHITKLNAENNAATPFKIDNVKKARVQV 275

OY 63 AGKKYFIDFVARETTCKESNEELTSCETKLGQSLDCNAEVYVVPWEKKIYPTV 118

DB 276 AGKKYFIDFVARETTCKESNEELTSCETKLGQSLDCNAEVYVVPWEKKIYPTV 331

RESULT 7

ABU99143

ID ABU99143 standard; protein; 398 AA.

AC ABU99143;

XX

XX 01-AUG-2003 (first entry)

XX Novel human GPCR related protein NOV12a.

XX Human; G-protein coupled receptor related protein; GPCR related protein;

XX NOV; cytotactic; cardiac; antiarteriosclerotic; antidiabetic;

XX immunomodulator; anti-HIV; anorectic; antisthmatic; haemostatic;

XX antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;

XX NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;

KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
 OS Homo sapiens.
 XX WO200299116-A2.
 XX PD 12-DEC-2002.
 XX PF 04-JUN-2002; 2002WO-US017428.
 XX PR 04-JUN-2001; 2001US-0295607P.
 XX PR 06-JUN-2001; 2001US-0295661P.
 XX PR 06-JUN-2001; 2001US-0296404P.
 XX PR 06-JUN-2001; 2001US-0296418P.
 XX PR 14-JUN-2001; 2001US-0298285P.
 XX PR 15-JUN-2001; 2001US-0298556P.
 XX PR 21-JUN-2001; 2001US-0299949P.
 XX PR 26-JUN-2001; 2001US-0300883P.
 XX PR 28-JUN-2001; 2001US-0301550P.
 XX PR 13-AUG-2001; 2001US-0311972P.
 XX PR 27-AUG-2001; 2001US-0315071P.
 XX PR 29-AUG-2001; 2001US-0315660P.
 XX PR 14-SEP-2001; 2001US-0322293P.
 XX PR 17-SEP-2001; 2001US-0327086P.
 XX PR 14-DEC-2001; 2001US-0341186P.
 XX PR 28-FEB-2002; 2002US-0361189P.
 XX PR 12-MAR-2002; 2002US-0363673P.
 XX PR 13-MAR-2002; 2002US-0363676P.
 XX PR 03-JUN-2002; 2002US-00363676.
 XX PA (CURA-) CURAGEN CORP.
 XX PI Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR,
 PI Gargolli EA, Gerlach VL, Gorman L, Guo X, Hjelt T, Kekuda R, Li L,
 PI Macdougall JR, Malyanar UM, Millet I, Padigaru M, Patturajan M,
 PI Pena CE, Rastelli L, Shinkens RA, Stone DJ, Spytek KA, Varnet CAM,
 PI Voss EZ, Zorhusen BB;
 XX WPI, 2003-140627/13.
 DR N-PSDB, ACD03647.
 XX

PT The invention describes an isolated polypeptide (I) comprising any of 27
 CC 118-961 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOVX-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC disorders, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis, in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOVX substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as

CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX Sequence 398 AA;
 SQ Query Match 90.0%; Score 618; DB 6; Length 398;
 Best Local Similarity 100.0%; Pred. No. 2.8e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVGCPRDPTNSPELETLTHITKLAENNAATYFKINDVVKARVQVV 62
 DB 224 GKDFVQPTKICVGCPRDPTNSPELETLTHITKLAENNAATYFKINDVVKARVQVV 283
 QY 63 AGKKYFIDFVARETTCKSNBELTESCETKLGSLDCNAEVVVPWEKKIYPTV 118
 DB 284 AGKKYFIDFVARETTCKSNBELTESCETKLGSLDCNAEVVVPWEKKIYPTV 339
 RESULT 8
 ADE76864
 ID ADE76864 standard; protein; 427 AA.
 XX AC ADE76864;
 XX DT 29-JAN-2004 (first entry)
 XX DE Human protein expressed in a liver disorder #9.
 XX KW human; liver disorder; hyperlipidaemia; hypertension; type II diabetes;
 KW tumour; liver; inflammatory disorder; immune response disorder;
 KW high-throughput screening; differential gene expression; gene therapy.
 XX OS Homo sapiens.
 XX FN US2003108871-A1.
 XX PD 12-JUN-2003.
 XX PF 30-JUL-2001; 2001US-00919039.
 XX PR 28-JUL-2000; 2000US-0222113P.
 XX PA (KASE/) KASER M R.
 XX PI Kaser MR;
 DR WPI; 2004-031227/03.
 DR N-PSDB; ADE76863.
 XX Composition comprising several cDNAs that are differentially expressed in
 PT treated human C3A liver cell cultures, useful for treating liver
 PT disorders.
 PS Claim 1; SEQ ID NO 29; 41pp; English.
 XX The invention relates to a composition comprising several cDNAs that are
 CC differentially expressed in a liver disorder. The composition is useful
 CC for treating liver disorder such as hyperlipidaemia, hypertension, type
 CC II diabetes, tumours of the liver and disorders of the inflammatory and
 CC immune response. The composition is useful for a high-throughput method
 CC of screening several molecules or compounds to identify a ligand which
 CC specifically binds a cDNA. A protein encoded by the cDNA is useful for a
 CC high-throughput method for using a protein to screen several molecules or
 CC compounds to identify at least one ligand which specifically binds the
 CC protein which involves combining the protein encoded by the cDNA with
 CC several of molecules or compounds under conditions to allow specific
 CC binding, and detecting specific binding between the protein and a
 CC molecule or compound, therefore identifying a ligand which specifically
 CC binds the protein. The composition is useful for detecting and
 CC quantifying differential gene expression, can be used in gene therapy, to
 CC formulate prognosis and to design a treatment regimen and to monitor the
 CC efficacy of treatment. The present sequence represents the amino acid
 CC sequence of a protein encoded by a cDNA differentially expressed in a

CC liver disorder.
 XX Sequence 427 AA;
 SQ

Query Match
 Best Local Similarity 90.0%; Score 618; DB 8; Length 427;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDVPQPTKICVCGPRDIPNTSPLEETLTHTITKLAENNAATFFPKIDNVKARVQV 62
 DB 253 GKDVPQPTKICVCGPRDIPNTSPLEETLTHTITKLAENNAATFFPKIDNVKARVQV 312

QY 63 AGKXYFIDFVARETTCKESNEELTSCETKLGQSLDCNAEVVVPWEKKIYPTV 118
 DB 313 AGKXYFIDFVARETTCKESNEELTSCETKLGQSLDCNAEVVVPWEKKIYPTV 368

RESULT 9
 ABU99144
 ID ABU99144 standard; protein: 615 AA.
 AC ABU99144;
 XX
 XX 01-AUG-2003 (first entry)
 XX Novel human GPCR related protein NOV12b.
 XX Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytosolic; cardiac; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antilasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nontropic; gene therapy; vaccine;
 KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
 XX Homo sapiens.
 OS
 XX
 XX WO200299116-A2.
 XX
 XX 12-DEC-2002.
 XX
 XX 04-JUN-2002; 2002WO-US017428.
 XX
 XX 04-JUN-2001; 2001US-0295607P.
 XX 04-JUN-2001; 2001US-0295661P.
 XX 06-JUN-2001; 2001US-0296404P.
 XX 06-JUN-2001; 2001US-0296418P.
 XX 14-JUN-2001; 2001US-0298285P.
 XX 15-JUN-2001; 2001US-0298556P.
 XX 21-JUN-2001; 2001US-0299549P.
 XX 26-JUN-2001; 2001US-0300833P.
 XX 28-JUN-2001; 2001US-0301550P.
 XX 13-AUG-2001; 2001US-0311972P.
 XX 27-AUG-2001; 2001US-0315071P.
 XX 29-AUG-2001; 2001US-0315660P.
 XX 14-SEP-2001; 2001US-0322933P.
 XX 17-SEP-2001; 2001US-0322706P.
 XX 14-DEC-2001; 2001US-0341186P.
 XX 28-FEB-2002; 2002US-0361189P.
 XX 12-MAR-2002; 2002US-0363673P.
 XX 12-MAR-2002; 2002US-0363676P.
 XX 03-JUN-2002; 2002US-00363676.
 XX (CURA-) CURAGEN CORP.
 XX
 XX Anderson DW, Baugartner JC, Boldog FB, Casman SJ, Edinger SR;
 PI Gangolli EA, Gerlach VL, Gorman L, Guo X, Hjalte T, Kekuda R, Li L;
 PI MacDougall JR, Malyankar DN, Millet I, Padigara M, Patturajan M;
 PI Pena CEA, Rastelli L, Shimkete RA, Stone SJ, Szytko KA, Vernet CAM;

PI Voss EZ, Zerhusen BD;
 XX WPI, 2003-140627/13.
 XX N-ESDS; ACDO3648.
 XX New NOVX polypeptides and nucleic acids, useful for preventing or
 PT treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,
 PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
 PT pharmacogenomics.
 XX
 XX Claim 1; Page 144; 332pp; English.
 XX The invention describes an isolated polypeptide (1) comprising any of 27
 CC 118-961 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOVX-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC disease, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infectious multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis, in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOVX substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridization probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX
 SQ Sequence 615 AA;
 Query Match 90.0%; Score 618; DB 6; Length 615;
 Best Local Similarity 100.0%; Pred. No. 5.18-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDVPQPTKICVCGPRDIPNTSPLEETLTHTITKLAENNAATFFPKIDNVKARVQV 62
 DB 224 GKDVPQPTKICVCGPRDIPNTSPLEETLTHTITKLAENNAATFFPKIDNVKARVQV 283

QY 63 AGKXYFIDFVARETTCKESNEELTSCETKLGQSLDCNAEVVVPWEKKIYPTV 118
 DB 284 AGKXYFIDFVARETTCKESNEELTSCETKLGQSLDCNAEVVVPWEKKIYPTV 339

RESULT 10
 ABB78707
 ID ABB78707 standard; protein: 626 AA.
 XX
 AC ABB78707;
 XX
 XX 18-JUL-2002 (first entry)
 XX Human high molecular weight kininogen (HK) mature protein SEQ ID NO:1.
 XX Human; kininogen; high molecular weight kininogen; HK; D5 domain;
 XX D5 receptor; angiogenesis; endothelial cell; cytostatic; antitumour;
 KW antithrombotic; vasotropic; vulnary; tranquilliser; thrombolytic;
 KW ophthalmological; gynaecological; antitumor; antidiabetic; antiarthritic;
 KW antiangiogenic; antiapoptotic; endocrine; apoptosis; gene therapy.
 XX Homo sapiens.
 OS
 XX
 XX Key Location/Qualifiers
 FH Domain 384..508
 FT /label= D5_domain
 XX
 XX WO200214369-A2.

XX PD 21-FEB-2002.
 XX XX
 XX PF 24-JUL-2001; 2001WO-US033185.
 XX XX
 XX PR 24-JUL-2000; 2000US-0220194P.
 XX XX
 XX PA (ATTE-) ATTENUON LLC.
 XX XX
 XX PI Mazar AP, Juarez JC;
 XX XX
 XX DR WPI; 2002-393611/42.
 XX XX
 XX PT Novel human kininogen D5 domain polypeptides useful for treating
 XX PT conditions associated with endothelial cell migration, proliferation,
 XX PT invasion or angiogenesis, e.g. arthritis, macular degeneration, benign
 XX PT hyperplasia.
 XX PS
 XX PS Disclosure; Page 13; 84pp; English.
 XX XX
 CC The present invention describes an isolated polypeptide (I) that
 CC corresponds to the D5 domain of human kininogen, or biologically active
 CC peptide fragment, homologue or functional derivative, and which: (a)
 CC inhibits angiogenesis; (b) binds to the D5 binding site on endothelial
 CC cells (EC); (c) activates signalling pathways leading to the introduction
 CC of apoptosis in EC; and/or (d) inhibits the signalling pathway required
 CC for maintenance of EC viability. (I) has cytostatic, antitumour,
 CC antiatherosclerotic, vasotropic, vulnerary, tranquiliser, thrombolytic,
 CC ophthalmological, gynaecological, antiulcer, antidiabetic, antiarthritic,
 CC antiangiogenic, antiapoptotic and endocrine activities. An antibody (IX)
 CC specific for an epitope of (I) is useful for inhibiting tumour growth or
 CC angiogenesis in a subject. (I), a D5 fusion polypeptide (II) or a dimeric
 CC or trimeric fusion polypeptide (III) can be used for inhibiting EC
 CC migration, proliferation, invasion, or angiogenesis, or for inducing EC
 CC apoptosis. An angiogenic EC-targeting pharmaceutical composition (X)
 CC comprising (I), (II), or (III), can be used for treating a subject having
 CC a disease or condition associated with undesired EC migration,
 CC proliferation, invasion or angiogenesis. (I), (II), or (III) can be used
 CC for isolating a D5 domain binding molecule from a complex mixture and for
 CC isolating or enriching cells expressing D5 domain binding sites from a
 CC cell mixture. The present sequence represents the mature human high
 CC molecular weight kininogen (HK) protein, which is given in the
 CC exemplification of the present invention
 XX SQ Sequence 626 AA;
 Query Match 90.0%; Score 618; DB 5; Length 626;
 Best Local Similarity 100.0%; Pred. No. 5.3e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 3 GKDVPPTKICVCGPRDPTNSPELETLTHITKLNANNATFYFKIDNVKARQVV 62
 Db 235 GKDVPPTKICVCGPRDPTNSPELETLTHITKLNANNATFYFKIDNVKARQVV 294
 Qy 63 AGKGYFIDFVARETTCKSKESNEELTESCTKGLQSLDCNNAEYVVPWEKKIYPTV 118
 Db 295 AGKGYFIDFVARETTCKSKESNEELTESCTKGLQSLDCNNAEYVVPWEKKIYPTV 350
 RESULT 11
 ABG21101
 ID ABG21101 standard; protein; 644 AA.
 AC ABG21101;
 XX ABG21101;
 XX 18-FEB-2002 (first entry)
 XX Novel human diagnostic protein #21092.
 DE Human; chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder.
 XX Homo sapiens.
 OS

XX PN WO200175067-A2.
 XX PD 11-OCT-2001.
 XX XX
 XX XX 30-MAR-2001; 2001WO-US008631.
 XX XX
 XX XX 31-MAR-2000; 2000US-00540217.
 XX PR 23-AUG-2000; 2000US-00649167.
 XX XX
 XX XX (HYSE-) HYSSEQ INC.
 XX XX
 XX XX Dmanac RT, Liu C, Tang YT;
 XX XX
 XX XX WPI; 2001-639162/73.
 XX DR N-PSDB; AS88288.
 XX XX
 XX PT New isolated polynucleotide and encoded polypeptides, useful in
 XX PT diagnostics, forensics, gene mapping, identification of mutations
 XX PT responsible for genetic disorders or other traits and to assess
 XX PT biodiversity.
 XX PS
 XX PS Claim 20; SEQ ID NO 51460; 103pp; English.
 XX XX
 CC The invention relates to isolated polynucleotide (I) and polypeptide (II)
 CC sequences. (I) is useful as hybridisation probes, polymerase chain
 CC reaction (PCR) primers, oligomers and for chromosome and gene mapping,
 CC and in recombinant production of (II). The polynucleotides are also used
 CC in diagnostics as expressed sequence tags for identifying expressed
 CC genes. (I) is useful in gene therapy techniques to restore normal
 CC activity of (II) or to treat disease states involving (II). (II) is
 CC useful for generating antibodies against it, detecting or quantitating a
 CC polypeptide in tissue, as molecular weight markers and as a food
 CC supplement. (II) and its binding partners are useful in medical imaging
 CC of sites expressing (II). (I) and (II) are useful for treating disorders
 CC involving aberrant protein expression or biological activities. The
 CC polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. ABG00010-ABG30377 represent novel human diagnostic
 CC amino acid sequences of the invention. Note: The sequence data for this
 CC patent did not appear in the printed specification, but was obtained in
 CC electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX SQ Sequence 644 AA;
 Query Match 90.0%; Score 618; DB 4; Length 644;
 Best Local Similarity 100.0%; Pred. No. 5.5e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 3 GKDVPPTKICVCGPRDPTNSPELETLTHITKLNANNATFYFKIDNVKARQVV 62
 Db 253 GKDVPPTKICVCGPRDPTNSPELETLTHITKLNANNATFYFKIDNVKARQVV 312
 Qy 63 AGKGYFIDFVARETTCKSKESNEELTESCTKGLQSLDCNNAEYVVPWEKKIYPTV 118
 Db 313 AGKGYFIDFVARETTCKSKESNEELTESCTKGLQSLDCNNAEYVVPWEKKIYPTV 368
 RESULT 12
 ABB78710
 ID ABB78710 standard; protein; 644 AA.
 XX ABB78710;
 AC ABB78710;
 XX ABB78710;
 XX 18-JUL-2002 (first entry)
 XX Human high molecular weight kininogen (HK) protein.
 DE Human; kininogen; high molecular weight kininogen; HK; D5 domain;
 KW D5 receptor; angiogenesis; endothelial cell; cytostatic; antitumour;
 KW

KW antiatherosclerotic; vasotropic; vulnerary; tranquilliser; thrombolytic;
 KW ophthalmological; gynaecological; antiulcer; antidiabetic; antiarthritic;
 KW antiangiogenic; antiapoptotic; endocrine; apoposis; gene therapy.
 XX
 OS Homo sapiens.

Key	Location/Qualifiers
Peptide	1..18
Protein	/label= signal
	19..644
	/label= mature_human_high_molecular_weight_kinogen
Disulfide-bond	28..614
Disulfide-bond	83..94
Disulfide-bond	107..126
Disulfide-bond	142..145
Disulfide-bond	206..218
Disulfide-bond	229..248
Disulfide-bond	264..267
Disulfide-bond	328..340
Disulfide-bond	351..370
Domain	402..526
	/label= D5_domain

WO200214369-A2.

21-FEB-2002.

24-JUL-2001; 2001WO-US023185.

24-JUL-2000; 2000US-0220194P.

(ATTE-) ATTENUON LLC.

Mazar AP, Juarez JC;

WPI, 2002-393611/42.

Novel human kinogen D5 domain polypeptides useful for treating conditions associated with endothelial cell migration, proliferation, invasion or angiogenesis, e.g. arthritis, macular degeneration, benign hyperplasia.

Disclosure: Fig 1B-E; 84pp; English.

The present invention describes an isolated polypeptide (I) that corresponds to the D5 domain of human kinogen, or biologically active peptide fragment, homologue or functional derivative, and which: (a) inhibits angiogenesis; (b) binds to the D5 binding site on endothelial cells (EC); (c) activates signalling pathways leading to the introduction of apoptosis in EC; and/or (d) inhibits the signalling pathway required for maintenance of EC viability. (I) has cytostatic, antitumour, antiatherosclerotic, vasotropic, vulnerary, tranquilliser, thrombolytic, ophthalmological, gynaecological, antiulcer, antidiabetic, antiarthritic, antiangiogenic, antiapoptotic and endocrine activities. An antibody (IX) specific for an epitope of (I) is useful for inhibiting tumour growth or angiogenesis in a subject. (II), a D5 fusion polypeptide (II) or a dimeric or trimeric fusion polypeptide (III) can be used for inhibiting EC migration, proliferation, invasion, or angiogenesis, or for inducing EC apoptosis. An angiogenic EC-targeting pharmaceutical composition (X) comprising (I), (II), or (III), can be used for treating a subject having a disease or condition associated with undesired EC migration, proliferation, invasion or angiogenesis. (I), (II), or (III) can be used for isolating a D5 domain binding molecule from a complex mixture and for isolating or enriching cells expressing D5 domain binding sites from a cell mixture. The present sequence represents the human high molecular weight kinogen (HK) protein, which is given in the exemplification of the present invention.

Sequence 644 AA;

Query Match 90.0%; Score 618; DB 5; Length 644;
 Best Local Similarity 100.0%; Pred. No. 5.5e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY	3	QKDFVQPTKICVCPDIDITNSPELBEFLTWITTKLAENNATPYFKIDNVKARQVV 62
DB	253	GKDFVQPTKICVCPDIDITNSPELBEFLTWITTKLAENNATPYFKIDNVKARQVV 312
QY	63	AGKYPIDFVARETTCKSENEELTSCETKXKLGSLDCNAEVVVPWEKKIYPTV 118
DB	313	AGKYPIDFVARETTCKSENEELTSCETKXKLGSLDCNAEVVVPWEKKIYPTV 368

RESULT 13

ABU99150
 ID ABU99150 standard; protein; 644 AA.

AC ABU99150;

XX 01-AUG-2003 (first entry)

DE Novel human GPCR related protein NOV12h.

KW Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytostatic; cardiant; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW antiparkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.

OS Homo sapiens.

FN WO2002299116-A2.

XX 12-DEC-2002.

PD 04-JUN-2002; 2002WO-US017428.

XX 04-JUN-2001; 2001US-0295607P.

PR 04-JUN-2001; 2001US-0235661P.

PR 06-JUN-2001; 2001US-0256404P.

PR 14-JUN-2001; 2001US-0298285P.

PR 15-JUN-2001; 2001US-0298556P.

PR 21-JUN-2001; 2001US-0299499P.

PR 26-JUN-2001; 2001US-0300883P.

PR 13-AUG-2001; 2001US-0311972P.

PR 27-AUG-2001; 2001US-0315071P.

PR 14-SEP-2001; 2001US-0322293P.

PR 17-SEP-2001; 2001US-0322706P.

PR 14-DEC-2001; 2001US-0341186P.

PR 28-FEB-2002; 2002US-0361189P.

PR 12-MAR-2002; 2002US-0363676P.

PR 03-JUN-2002; 2002US-00363676.

(CURA-) CURAGEN CORP.

PA Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR;
 PI Ganggoli EA, Gerlach VL, Gorman L, Guo X, Hjalte T, Kekuda R, Li L;
 PI Macdougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M;
 PI Pena CEA, Rastelli L, Shinketa RA, Stone DJ, Spytek KA, Vernet CM;
 PI Voss EZ, Zerhusen BD;
 XX WPI, 2003-140627/13.
 DR N-PSDB; AC003654.

XX New NOVX polypeptides and nucleic acids, useful for preventing or
 PI treating NOVX-associated disorders, e.g. cancer, cardiomyopathy,

PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
 PT pharmacogenomics.
 XX Claim 1, Page 148; 332pp; English.
 XX
 CC The invention describes an isolated polypeptide (I) comprising any of 27
 CC 118-961 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOVX-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC diseases, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis, in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOVX substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX Sequence 644 AA;
 SQ
 Query Match 90.0%; Score 618; DB 6; Length 644;
 Best Local Similarity 100.0%; Pred. No. 5.5e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVCGPRDIPNPSPELEETLTHITKLNANNATFYFKIDNVKARQV 62
 DB 253 GKDFVQPTKICVCGPRDIPNPSPELEETLTHITKLNANNATFYFKIDNVKARQV 312
 QY 63 AGKYTFIDFVARETTCSKESNEELTESCETKKGQSLDCNAEYVVPPEKKIYPTV 118
 DB 313 AGKYTFIDFVARETTCSKESNEELTESCETKKGQSLDCNAEYVVPPEKKIYPTV 368
 RESULT 14
 ABU99145
 ID ABU99145 standard; protein; 644 AA.
 AC ABU99145;
 XX
 DT 01-AUG-2003 (first entry)
 DE Novel human GPCR related protein NOV12c.
 KW Human; G-protein coupled receptor related protein; GPCR related protein;
 KW NOV; cytosolic; cardiac; antiarteriosclerotic; antidiabetic;
 KW immunomodulator; anti-HIV; anorectic; antiasthmatic; haemostatic;
 KW anti-parkinsonian; neuroprotective; nootropic; gene therapy; vaccine;
 KW NOVX-associated disorder; cardiomyopathy; atherosclerosis; cancer;
 KW diabetes; immune disorder; AIDS; obesity; asthma;
 KW haematopoietic disorder; Parkinson's disease; Alzheimer's disease;
 KW infection; multiple sclerosis; cancer-associated cachexia;
 KW wasting disorder; chronic disease; neurogenesis; cell differentiation;
 KW cell proliferation; haematopoiesis; wound healing; angiogenesis;
 KW chromosome mapping; tissue typing; preventive medicine; pharmacogenomic.
 XX Homo sapiens.
 OS
 XX WO200299116-A2.
 PN
 XX
 PD 12-DEC-2002.
 XX
 PP 04-JUN-2002; 2002WO-US017428.
 XX

PR 04-JUN-2001; 2001US-0295607P.
 PR 04-JUN-2001; 2001US-0295661P.
 PR 06-JUN-2001; 2001US-0296404P.
 PR 06-JUN-2001; 2001US-0296418P.
 PR 14-JUN-2001; 2001US-0298285P.
 PR 14-JUN-2001; 2001US-0298285P.
 PR 18-JUN-2001; 2001US-0298556P.
 PR 21-JUN-2001; 2001US-0299949P.
 PR 28-JUN-2001; 2001US-0300883P.
 PR 28-JUN-2001; 2001US-0301550P.
 PR 13-AUG-2001; 2001US-0311972P.
 PR 27-AUG-2001; 2001US-0315071P.
 PR 29-AUG-2001; 2001US-0315660P.
 PR 14-SEP-2001; 2001US-032293P.
 PR 17-SEP-2001; 2001US-0322706P.
 PR 14-DEC-2001; 2001US-0341186P.
 PR 28-FEB-2002; 2002US-0361189P.
 PR 12-MAR-2002; 2002US-0363673P.
 PR 12-MAR-2002; 2002US-0363676P.
 PR 03-JUN-2002; 2002US-0363676.
 XX
 PA (CURA-) CUPAGEN CORP.
 XX
 PI Anderson DW, Baumgartner JC, Boldog FL, Casman SJ, Edinger SR,
 PI Gangoli EA, Gerlach VL, Gorman L, Guo X, Hjalit T, Kekuda R, Li L,
 PI McDougall JR, Malyankar UM, Millet I, Padigaru M, Patturajan M,
 PI Pena CE, Rastelli L, Shinkets RA, Stone DJ, Spytek KA, Vernet CAM,
 PI Voss EZ, Zerhusen BD;
 XX WPI; 2003-140627/13.
 DR N-PSDB; ACD03649.
 XX
 PT New NOVX polypeptides and nucleic acids, useful for preventing or
 PT treating NOVX-associated disorders, e.g. cancer, cardiomyopathy, or
 PT atherosclerosis, or diabetes, and in chromosome mapping, tissue typing or
 PT pharmacogenomics.
 XX
 FS Claim 1; Page 144-145; 332pp; English.
 XX
 CC The invention describes an isolated polypeptide (I) comprising any of 27
 CC 118-961 residue amino acid sequences, given in the specification, a
 CC mature form of them, a sequence that is at least 95 % identical to them,
 CC or a sequence having one or more conservative substitutions in them. The
 CC polypeptide is useful in manufacturing a medicament for treating a
 CC syndrome associated with a human disease selected from a pathology
 CC associated with the polypeptide. The NOVX polypeptides, polynucleotides
 CC and antibodies are useful in treating or preventing NOVX-associated
 CC disorders, e.g. cardiomyopathy, atherosclerosis, cancer, diabetes, immune
 CC diseases, AIDS, obesity, asthma, haematopoietic disorders, Parkinson's
 CC disease, Alzheimer's disease, infections, multiple sclerosis, cancer-
 CC associated cachexia, and other wasting disorders associated with chronic
 CC diseases. The nucleic acids and polypeptides may also be used as targets
 CC for the identification of small molecules that modulate or inhibit e.g.
 CC neurogenesis, cell differentiation, cell proliferation, haematopoiesis,
 CC wound healing and angiogenesis, in gene therapy, in generation of
 CC antibodies that bind immunospecifically to NOVX substances for use in
 CC therapeutic or diagnostic methods. The nucleic acids are further used as
 CC hybridisation probes, in chromosome mapping, tissue typing, preventive
 CC medicine, and pharmacogenomics. The polypeptides are also useful as
 CC vaccines. This is the amino acid sequence of a novel human G-protein
 CC coupled receptor related protein NOV
 XX Sequence 644 AA;
 SQ
 Query Match 90.0%; Score 618; DB 6; Length 644;
 Best Local Similarity 100.0%; Pred. No. 5.5e-62;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVCGPRDIPNPSPELEETLTHITKLNANNATFYFKIDNVKARQV 62
 DB 253 GKDFVQPTKICVCGPRDIPNPSPELEETLTHITKLNANNATFYFKIDNVKARQV 312
 QY 63 AGKYTFIDFVARETTCSKESNEELTESCETKKGQSLDCNAEYVVPPEKKIYPTV 118

Db 313 AGKXFDVARETTCSENEBELTESCETKXQSLDCNAEVVYVPWEKKIYPTV 368

RESULT 15

AAB37447

ID AAB37447 standard; protein; 122 AA.

XX AC AAB37447;

XX DT 21-FEB-2001 (first entry)

XX DE Human kininogen D3.

XX KW Enzyme; legumain; endopeptidase; Cystatin; human; kininogen.

XX OS Homo sapiens.

XX PN WO200064945-A1.

XX PD 02-NOV-2000.

XX PP 20-APR-2000; 2000WO-GB001571.

XX PR 22-APR-1999; 99GB-00009133.

XX PA (BADR-) BABRAHAM INST.

XX PI Abrahamson M, Barrett AJ;

XX DR WPI; 2000-687316/67.

XX PT Inhibition of mammalian legumain or legumain-related endopeptidase by
 PT cystatin involves interaction with second papain-non-reactive site of
 PT cystatin.

XX PS Disclosure; Fig 4; 45pp; English.

XX CC The present invention relates to inhibition of the enzymatic activity of
 CC legumain or a legumain-related endopeptidase by cystatin. The inhibition
 CC involves an interaction between legumain and a papain-non-reactive site
 CC of cystatin. Legumain (EC 3.4.22.34) is a cysteine endopeptidase, and
 CC performs a protein-processing function. The present sequence is human
 CC kininogen D3, which was used in the present invention. Kininogen is a
 CC type 3 cystatin

XX SQ Sequence 122 AA;

Query Match 85.3%; Score 586; DB 3; Length 122;

Best Local Similarity 100.0%; Pred. No. 2-7e-59;

Matches 110; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 9 PPTKICVGCPRDPTSPLEBELTHTITKLAENNAATFYKIDNVKARVQVWAGKKYP 68

DB 1 PPTKICVGCPRDPTSPLEBELTHTITKLAENNAATFYKIDNVKARVQVWAGKKYP 60

OY 69 IDPVARETTCSENEBELTESCETKXQSLDCNAEVVYVPWEKKIYPTV 118

DB 61 IDPVARETTCSENEBELTESCETKXQSLDCNAEVVYVPWEKKIYPTV 110

Search completed: September 24, 2004, 14:08:38

Job time : 52.308 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 24, 2004, 14:07:01 / Search time 14.732 Seconds
(without alignments)
445.051 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 687

Sequence: 1 GSKGFVOPPKICVGCPRD.....VPWEKKIYPTTVVHNECEP 127

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 389414 seqs, 51625971 residues

Total number of hits satisfying chosen parameters: 389414

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued Patents AA.*
1: /cgn2_6/ptodata/2/iaa/5A COMB.pcp.*
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3: /cgn2_6/ptodata/2/iaa/6A COMB.pcp.*
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5: /cgn2_6/ptodata/2/iaa/PCUTS COMB.pcp.*
6: /cgn2_6/ptodata/2/iaa/backfiles1.pcp.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	558	81.2	117	1	US-08-193-114B-1
2	556.5	81.0	117	5	PCT-US92-06809-1
3	163.5	23.8	145	2	US-08-832-535-2
4	163.5	23.8	145	3	US-09-019-485-2
5	163.5	23.8	145	3	US-09-019-485-3
6	163.5	23.8	145	3	US-09-431-480-9
7	163.5	23.8	145	3	US-09-617-302-9
8	163.5	23.8	145	4	US-09-528-4368-2
9	163	23.7	178	2	US-08-791-522-1
10	163	23.7	178	3	US-09-314-777-1
11	138.5	20.2	121	4	US-09-775-932-14
12	138.5	20.2	128	4	US-09-775-932-12
13	138.5	20.2	149	2	US-08-461-030C-2
14	138.5	20.2	149	3	US-08-744-138-2
15	138.5	20.2	149	3	US-09-431-480-8
16	138.5	20.2	149	3	US-09-431-480-10
17	138.5	20.2	149	3	US-09-617-302-8
18	138.5	20.2	149	3	US-09-617-302-10
19	138.5	20.2	149	4	US-09-241-376-3
20	138.5	20.2	149	4	US-09-940-497-2
21	137.5	20.0	112	4	US-08-843-303-16
22	136.5	19.9	118	4	US-09-775-932-24
23	135.5	19.7	146	6	5432264-6
24	134	19.5	148	5	PCT-US95-07135-2
25	132.5	19.3	120	4	US-09-775-932-2
26	132.5	19.3	145	2	US-08-832-535-11
27	132.5	19.3	146	2	US-08-791-522-3

28	132.5	19.3	146	3	US-08-744-138-3
29	132.5	19.3	146	3	US-09-019-485-4
30	132.5	19.3	146	3	US-09-314-777-3
31	132.5	19.3	146	3	US-09-431-480-6
32	132.5	19.3	146	3	US-09-617-302-6
33	132.5	19.3	146	4	US-09-241-376-3
34	132.5	19.3	146	4	US-09-528-4368-3
35	132.5	19.3	146	4	US-09-886-319A-47
36	132.5	19.3	146	4	US-09-940-497-3
37	132.5	19.3	146	4	US-09-976-594-37
38	132.5	19.3	146	4	US-08-849-303-17
39	132.5	19.3	146	5	PCT-US95-07135-9
40	132	19.2	26	3	US-08-676-242-15
41	131.5	19.1	382	4	US-09-599-1608-93
42	130	18.9	127	4	US-08-849-303-13
43	129.5	18.9	140	4	US-09-886-319A-46
44	129.5	18.9	140	4	US-09-886-319A-48
45	128	18.6	111	4	US-08-849-303-26

ALIGNMENTS

RESULT 1

US-08-193-114B-1

Sequence 1, Application US/08193114B

Patent No. 5472945

GENERAL INFORMATION:

APPLICANT: Schmaier, Alvin H.

APPLICANT: Jiang, Yongping

TITLE OF INVENTION: Modulation of Blood

TITLE OF INVENTION: Pressure and Inhibition of Platelet Activation

TITLE OF INVENTION: with Kininogen Fragment

NUMBER OF SEQUENCES: 2

CORRESPONDENCE ADDRESS:

ADDRESSEE: Seidel, Gonda, Lavorgna &

ADDRESSEE: Monaco, P.C.

STREET: 1800 Two Penn Center Plaza

CITY: Philadelphia

STATE: Pennsylvania

COUNTRY: U.S.A.

ZIP: 19102

COMPUTER READABLE FORM: Diskette, 3.50 inch, 720 Kb

MEDIUM TYPE: IBM PS/2

COMPUTER: IBM PS/2

OPERATING SYSTEM: MS-DOS

SOFTWARE: WordPerfect 5.1

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/193,114B

FILING DATE: 9 February 1994

CLASSIFICATION: 514

PRIOR APPLICATION DATA:

APPLICATION NUMBER: U.S. Application

APPLICATION NUMBER: Serial No. 5472945 07/744,545

FILING DATE: 13 August 1991

ATTORNEY/AGENT INFORMATION:

NAME: Monaco, Daniel A.

REGISTRATION NUMBER: 30,480

REFERENCE/DOCKET NUMBER: 6056-137 CII

TELECOMMUNICATION INFORMATION:

TELEPHONE: (215) 568-8383

TELEFAX: (215) 568-5549

TELEX: No. 5472945e

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 117 amino acids

TYPE: peptide

TOPOLOGY: linear

US-08-193-114B-1

Query Match 81.2%; Score 558; DB 1; Length 117;
Best Local Similarity 100.0%; Pred. No. 2,3e-55;
Matches 105; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 14 CVGCPDIPNTPNSPELBTHTITKLNAENNATFYKIDNVKARQVWAGKKYPIDFVA 73
 DB 1 CVGCPDIPNTPNSPELBTHTITKLNAENNATFYKIDNVKARQVWAGKKYPIDFVA 60
 OY 74 RETTCSKESNEELTESCETKKGSLDCNAEVVVPWEKKIYPTV 118
 DB 61 RETTCSKESNEELTESCETKKGSLDCNAEVVVPWEKKIYPTV 105

RESULT 2
 PCT-US92-06809-1
 ; Sequence 1, Application PC/TUS9206809
 ; GENERAL INFORMATION:
 ; APPLICANT: Schmaier, Alvin H.
 ; APPLICANT: Jiang, Yongping
 ; TITLE OF INVENTION: Modulation of Blood
 ; TITLE OF INVENTION: Pressure by Altering Bradykinin Levels
 ; NUMBER OF SEQUENCES: 2
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Temple University - Of the
 ; ADDRESSEE: Commonwealth System of Higher Education
 ; STREET: 406 University Services
 ; CITY: Philadelphia
 ; STATE: Pennsylvania
 ; COUNTRY: U.S.A.
 ; ZIP: 19122

COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette, 3.50 inch, 720 Kb
 COMPUTER: IBM PS/2
 OPERATING SYSTEM: MS-DOS
 SOFTWARE: WordPerfect 5.1
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: PCT/US92/06809
 FILING DATE: 19910813
 CLASSIFICATION: 514
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: U.S. Application
 APPLICATION NUMBER: Serial No. 744,545
 FILING DATE: 13 August 1991
 ATTORNEY/AGENT INFORMATION:
 NAME: Monaco, Daniel A.
 REGISTRATION NUMBER: 30,480
 REFERENCE/DOCKET NUMBER: 6056-137
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (215) 568-8383
 TELEFAX: (215) 568-5549
 TELEX:
 INFORMATION FOR SEQ ID NO: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 117 amino acids
 TYPE: AMINO ACID
 TOPOLOGY: linear

PCT-US92-06809-1
 Query Match 81.0%; Score 556.5; DB 5; Length 117;
 Best Local Similarity 93.8%; Pred. No. 3.3e-55;
 Matches 106; Conservative 1; Mismatches 1; Indels 5; Gaps 1;
 OY 14 CVGCPDIPNTPNSPELBTHTITKLNAENNATFYKIDNVKARQVWAGKKYPIDFVA 73
 DB 1 CVGCPDIPNTPNSPELBTHTITKLNAENNATFYKIDNVKARQVWAGKKYPIDFVA 60
 OY 74 RETTCSKESNEELTESCETKKGSLDCNAEVVVPWEKKIYPTV 126
 DB 61 RETTCSKESNEELTESCETKKGSLDCNAEVVVPWEKKIYPTV 108

RESULT 3
 US-08-832-535-2
 ; Sequence 2, Application US/08812535
 ; Patent No. 5919658

GENERAL INFORMATION:
 APPLICANT: NI, JIAN
 APPLICANT: LI, HAODONG
 APPLICANT: YU, GUO-LIANG
 APPLICANT: GENTZ, REINER L.
 TITLE OF INVENTION: HUMAN CYSTATIN F
 NUMBER OF SEQUENCES: 11
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: HUMAN GENOME SCIENCES, INC.
 STREET: 9410 KEY WEST AVENUE
 CITY: ROCKVILLE
 STATE: MD
 COUNTRY: US
 ZIP: 20850
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/832,535
 FILING DATE: 03-APR-1997
 CLASSIFICATION: 435
 ATTORNEY/AGENT INFORMATION:
 NAME: KIMBALL, PAUL C.
 REGISTRATION NUMBER: 34,610
 REFERENCE/DOCKET NUMBER: PP265
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (201) 994-1700
 TELEFAX: (201) 994-1744
 INFORMATION FOR SEQ ID NO: 2:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 145 amino acids
 TYPE: amino acid
 TOPOLOGY: linear
 MOLECULE TYPE: Protein
 US-08-832-535-2

Query Match 23.8%; Score 163.5; DB 2; Length 145;
 Best Local Similarity 31.6%; Pred. No. 9.4e-11;
 Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;
 OY 11 TKICVGCPRDIPNTPNSPELBTHTITKLNAENNATFYKIDNVKARQVWAGKKYPIDF 70
 DB 32 SRVKGPPPTKTNDPGVLAARYSEKFNCTNDMPFKESRITRALVQVGLKTMLE 91
 OY 71 FVARETTCKSKSNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVTVNHWE 124
 DB 92 VEIGRTTCKGQHLRL-DDCDFTQHTLTKQTLSCYSEVWVVPW-----LQHEE 138

RESULT 4
 US-09-019-485-2
 ; Sequence 2, Application US/09019485
 ; Patent No. 6066617
 ; GENERAL INFORMATION:
 ; APPLICANT: Li, Haodong
 ; APPLICANT: Yu, Guo-Liang
 ; APPLICANT: Gentz, Reiner
 ; APPLICANT: NI, Jian
 ; TITLE OF INVENTION: Cystatin F
 ; NUMBER OF SEQUENCES: 17
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Human Genome Sciences, Inc.
 ; STREET: 9410 Key West Avenue
 ; CITY: Rockville
 ; STATE: MD
 ; COUNTRY: US
 ; ZIP: 20850
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS

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; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/019,485
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Benson, Robert H.
; REGISTRATION NUMBER: 30,446
; REFERENCE/DOCKET NUMBER: PP265P1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 3013098504
; TELEFAX: 3013098439
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 145 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-09-019-485-2

Query Match      23.8%; Score 163.5; DB 3; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.4e-11;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

Qy 11 TKICVCGPRDPTNSPELSELTHTITTKLNAENNATFYFKIDNVKARVQVAGKKYFID 70
Db 32 SRVKGPPKTIKTNDPGVLOAARYSVEKFNCTNDMFLFKESRITRALVQIVKGLKYMLE 91
Qy 71 FVARETTCSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVTNNHWE 124
Db 92 VEIGRTTCKKQNHRL--DDCDPQTNTLTKQLTSLCYSEVVVWVFW-----LQHFE 138

RESULT 5
US-09-019-485-3
; Sequence 3, Application US/09019485
; Patent No. 6086617
; GENERAL INFORMATION:
; APPLICANT: Li, Haodong
; APPLICANT: Yu, Guo-Liang
; APPLICANT: Gentz, Reiner
; APPLICANT: Ni, Jian
; TITLE OF INVENTION: Cystatin F
; NUMBER OF SEQUENCES: 17
; CORRESPONDENCE ADDRESS:
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: MD
; COUNTRY: US
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/019,485
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Benson, Robert H.
; REGISTRATION NUMBER: 30,446
; REFERENCE/DOCKET NUMBER: PP265P1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 3013098504
; TELEFAX: 3013098439
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 145 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-09-019-485-3

Query Match      23.8%; Score 163.5; DB 3; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.4e-11;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

Qy 11 TKICVCGPRDPTNSPELSELTHTITTKLNAENNATFYFKIDNVKARVQVAGKKYFID 70
Db 32 SRVKGPPKTIKTNDPGVLOAARYSVEKFNCTNDMFLFKESRITRALVQIVKGLKYMLE 91
Qy 71 FVARETTCSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVTNNHWE 124
Db 92 VEIGRTTCKKQNHRL--DDCDPQTNTLTKQLTSLCYSEVVVWVFW-----LQHFE 138

RESULT 6
US-09-431-480-9
; Sequence 9, Application US/09411480
; Patent No. 6235708
; GENERAL INFORMATION:
; APPLICANT: Holloway, James L.
; APPLICANT: Feldhaus, Andrew
; TITLE OF INVENTION: TESTIS SPECIFIC CYSTATIN-LIKE PROTEIN CYSTATIN T
; FILE REFERENCE: 98-72
; CURRENT APPLICATION NUMBER: US/09/431,480
; CURRENT FILING DATE: 1999-11-01
; EARLIER APPLICATION NUMBER: 60/109,217
; EARLIER FILING DATE: 1998-11-20
; EARLIER APPLICATION NUMBER: 60/156,382
; EARLIER FILING DATE: 1999-09-28
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 9
; LENGTH: 145
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-431-480-9

Query Match      23.8%; Score 163.5; DB 3; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.4e-11;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

Qy 11 TKICVCGPRDPTNSPELSELTHTITTKLNAENNATFYFKIDNVKARVQVAGKKYFID 70
Db 32 SRVKGPPKTIKTNDPGVLOAARYSVEKFNCTNDMFLFKESRITRALVQIVKGLKYMLE 91
Qy 71 FVARETTCSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVTNNHWE 124
Db 92 VEIGRTTCKKQNHRL--DDCDPQTNTLTKQLTSLCYSEVVVWVFW-----LQHFE 138

RESULT 7
US-09-617-302-9
; Sequence 9, Application US/09617302
; Patent No. 6245529
; GENERAL INFORMATION:
; APPLICANT: Holloway, James L.
; APPLICANT: Feldhaus, Andrew
; TITLE OF INVENTION: TESTIS SPECIFIC CYSTATIN-LIKE PROTEIN CYSTATIN T
; FILE REFERENCE: 98-72 Cl
; CURRENT APPLICATION NUMBER: US/09/617,302
; CURRENT FILING DATE: 2000-07-17
; PRIOR APPLICATION NUMBER: 09/431,480
; PRIOR FILING DATE: 1999-11-01
; PRIOR APPLICATION NUMBER: 60/109,217
; PRIOR FILING DATE: 1998-11-20
; PRIOR APPLICATION NUMBER: 60/156,382
; PRIOR FILING DATE: 1999-09-28
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 9
; LENGTH: 145
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; MOLECULE TYPE: protein
; US-09-019-485-3

Query Match      23.8%; Score 163.5; DB 3; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.4e-11;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

Qy 11 TKICVCGPRDPTNSPELSELTHTITTKLNAENNATFYFKIDNVKARVQVAGKKYFID 70
Db 32 SRVKGPPKTIKTNDPGVLOAARYSVEKFNCTNDMFLFKESRITRALVQIVKGLKYMLE 91
Qy 71 FVARETTCSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVTNNHWE 124
Db 92 VEIGRTTCKKQNHRL--DDCDPQTNTLTKQLTSLCYSEVVVWVFW-----LQHFE 138

RESULT 6
US-09-431-480-9
; Sequence 9, Application US/09411480
; Patent No. 6235708
; GENERAL INFORMATION:
; APPLICANT: Holloway, James L.
; APPLICANT: Feldhaus, Andrew
; TITLE OF INVENTION: TESTIS SPECIFIC CYSTATIN-LIKE PROTEIN CYSTATIN T
; FILE REFERENCE: 98-72
; CURRENT APPLICATION NUMBER: US/09/431,480
; CURRENT FILING DATE: 1999-11-01
; EARLIER APPLICATION NUMBER: 60/109,217
; EARLIER FILING DATE: 1998-11-20
; EARLIER APPLICATION NUMBER: 60/156,382
; EARLIER FILING DATE: 1999-09-28
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 9
; LENGTH: 145
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-431-480-9

Query Match      23.8%; Score 163.5; DB 3; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.4e-11;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

Qy 11 TKICVCGPRDPTNSPELSELTHTITTKLNAENNATFYFKIDNVKARVQVAGKKYFID 70
Db 32 SRVKGPPKTIKTNDPGVLOAARYSVEKFNCTNDMFLFKESRITRALVQIVKGLKYMLE 91
Qy 71 FVARETTCSKESNEELTESCE---TKKLGSLDCNAEVVVPWEKKIYPTVTNNHWE 124
Db 92 VEIGRTTCKKQNHRL--DDCDPQTNTLTKQLTSLCYSEVVVWVFW-----LQHFE 138

RESULT 7
US-09-617-302-9
; Sequence 9, Application US/09617302
; Patent No. 6245529
; GENERAL INFORMATION:
; APPLICANT: Holloway, James L.
; APPLICANT: Feldhaus, Andrew
; TITLE OF INVENTION: TESTIS SPECIFIC CYSTATIN-LIKE PROTEIN CYSTATIN T
; FILE REFERENCE: 98-72 Cl
; CURRENT APPLICATION NUMBER: US/09/617,302
; CURRENT FILING DATE: 2000-07-17
; PRIOR APPLICATION NUMBER: 09/431,480
; PRIOR FILING DATE: 1999-11-01
; PRIOR APPLICATION NUMBER: 60/109,217
; PRIOR FILING DATE: 1998-11-20
; PRIOR APPLICATION NUMBER: 60/156,382
; PRIOR FILING DATE: 1999-09-28
; NUMBER OF SEQ ID NOS: 22
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 9
; LENGTH: 145
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/ TYPE: PRT
/ ORGANISM: Homo sapiens
US-09-617-302-9

Query Match      23.8%; Score 163.5; DB 3; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.4e-11;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

QY 11 TKICVCPDIPNTPSPELETLTHITKLNAENNAATFYKIDNVKARVQVAGKYFID 70
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 32 SRVXGPPKTIKNDPGVLOAARYSVKFNCTNDMFLPKESRITRALVQIVKGLKYLE 91
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
QY 71 FVARETTCKESNEELTESCE---TKQLQSLDCNAEYVVPWEKKIYPTVTNHW 124
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 92 VEIGRTTCKNQHLRL-DDCDFQTNHTLKTLSYSEVWVVPW-----LQHP 138
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :

RESULT 8
US-09-528-436B-2
/ Sequence 2, Application US/09528436B
/ Patent No. 6576745
/ GENERAL INFORMATION:
/ APPLICANT: LI, et al.
/ TITLE OF INVENTION: Human Cystatin P
/ FILE REFERENCE: PF265PDI1
/ CURRENT APPLICATION NUMBER: US/09/528,436B
/ PRIOR FILING DATE: 2000-03-17
/ PRIOR APPLICATION NUMBER: 09/019,485
/ PRIOR FILING DATE: 1998-01-29
/ PRIOR APPLICATION NUMBER: 08/832,535
/ PRIOR FILING DATE: 1999-04-03
/ PRIOR APPLICATION NUMBER: 60/014,795
/ PRIOR FILING DATE: 1996-04-03
/ NUMBER OF SEQ ID NOS: 16
/ SOFTWARE: Patent in version 3.2
/ SEQ ID NO 2
/ LENGTH: 145
/ TYPE: PRT
/ ORGANISM: Homo sapiens
US-09-528-436B-2

Query Match      23.8%; Score 163.5; DB 4; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.4e-11;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

QY 11 TKICVCPDIPNTPSPELETLTHITKLNAENNAATFYKIDNVKARVQVAGKYFID 70
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 32 SRVXGPPKTIKNDPGVLOAARYSVKFNCTNDMFLPKESRITRALVQIVKGLKYLE 91
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
QY 71 FVARETTCKESNEELTESCE---TKQLQSLDCNAEYVVPWEKKIYPTVTNHW 124
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db 92 VEIGRTTCKNQHLRL-DDCDFQTNHTLKTLSYSEVWVVPW-----LQHP 138
   : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :

RESULT 9
US-08-791-522-1
/ Sequence 1, Application US/08791522
/ Patent No. 5935817
/ GENERAL INFORMATION:
/ APPLICANT: Bandman, Olga
/ APPLICANT: Goli, Suzya K.
/ TITLE OF INVENTION: NOVEL HUMAN CYSTATIN-LIKE
/ TITLE OF INVENTION: PROTEIN
/ NUMBER OF SEQUENCES: 4
/ CORRESPONDENCE ADDRESS:
/ ADDRESSEE: Incyte Pharmaceuticals, Inc.
/ STREET: 3174 Porter Drive
/ CITY: Palo Alto
/ STATE: CA
/ COUNTRY: USA
/ ZIP: 94304
/ COMPUTER READABLE FORM:
/ MEDIUM TYPE: Diskette
/ OPERATING SYSTEM: DOS
/ SOFTWARE: FastSeq for Windows Version 2.0
/ CURRENT APPLICATION DATA: US/09/314,777
/ FILING DATE:
/ CLASSIFICATION:
/ PRIOR APPLICATION DATA:
/ PRIOR APPLICATION NUMBER: 08/791,522
/ FILING DATE:
/ ATTORNEY/AGENT INFORMATION:
/ NAME: Billings, Lucy J.
/ REGISTRATION NUMBER: 36,749
/ REFERENCE/DOCKET NUMBER: PF-0193 US
```

TELECOMMUNICATION INFORMATION:

TELEPHONE: 415-855-0555
 TELEFAX: 415-845-4166
 INFORMATION FOR SEQ ID NO: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 178 amino acids
 TYPE: amino acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 IMMEDIATE SOURCE:
 CLONE: 30443
 US-09-314-777-1

Query Match 23.7%, Score 163, DB 3, Length 178,
 Best Local Similarity 34.0%, Pred. No. 1.4e-10,
 Matches 35, Conservative 20, Mismatches 44, Indels 4, Gaps 2;

Qy 11 TKICVCGPRDIPNPSPELEETLTHITKLNAENNATPYFKIDNVKARVQVAGKKYPI 70
 Db 54 SRVKGPFPTIKTNDPGVLOARYSVEKFNCTNDMPFKESRIITRALVOIVKGLKTYLE 113
 Qy 71 FVARETTCSKESENEELTESCE---TKLGGSLDCNAEVVVPW 110
 Db 114 VELGRITCKKNQHLRL-DDCDFOINHTLKLTLSCYSEVWVWP 155

RESULT 11

US-09-775-932-14;
 Sequence 14, Application US/09775932

Patent No. 6534477

GENERAL INFORMATION:

APPLICANT: University of British Columbia

TITLE OF INVENTION: Production and use of Modified Cystatins

FILE REFERENCE: 58069

CURRENT APPLICATION NUMBER: US/09/775.932

PRIOR FILING DATE: 2001-02-02

PRIOR APPLICATION NUMBER: CA99/00717

PRIOR FILING DATE: 1999-08-05

PRIOR APPLICATION NUMBER: 60/095,503

PRIOR FILING DATE: 1998-08-05

SOFTWARE: Patent in Ver. 2.0

SEQ ID NO 14

LENGTH: 121

TYPE: PRT

ORGANISM: Homo sapiens

US-09-775-932-14

Query Match

Best Local Similarity 20.2%, Score 138.5, DB 4, Length 121,
 Matches 34, Conservative 22, Mismatches 45, Indels 7, Gaps 3;

Qy 10 PTKICVCGPRDIPNPSPELEETLTHITKLNAENNATPYFKIDNVKARVQVAGKKYPI 69
 Db 2 PGRMVGELRLDSPDPQVOKAAQAAVASYNMGSNIYFRDTHIIKAQSLVAGIKYFL 61

Qy 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111

Db 62 TMEMGSTDCRTRVGTGDHVDLT-TCPLAAGAQOEKLCDFEVLVVPWQ 108

RESULT 12

US-09-775-932-12

Sequence 12, Application US/09775932

Patent No. 6534477

GENERAL INFORMATION:

APPLICANT: University of British Columbia

TITLE OF INVENTION: Production and use of Modified Cystatins

FILE REFERENCE: 58069

CURRENT APPLICATION NUMBER: US/09/775.932

PRIOR FILING DATE: 2001-02-02

PRIOR APPLICATION NUMBER: CA99/00717

PRIOR FILING DATE: 1999-08-05

PRIOR APPLICATION NUMBER: 60/095,503

PRIOR FILING DATE: 1998-08-05

NUMBER OF SEQ ID NOS: 32

SOFTWARE: Patent in Ver. 2.0

SEQ ID NO 12

LENGTH: 128

TYPE: PRT

ORGANISM: Homo sapiens

US-09-775-932-12

Query Match

Best Local Similarity 20.2%, Score 138.5, DB 4, Length 128,
 Matches 34, Conservative 22, Mismatches 45, Indels 7, Gaps 3;

Qy 10 PTKICVCGPRDIPNPSPELEETLTHITKLNAENNATPYFKIDNVKARVQVAGKKYPI 69
 Db 9 PGRMVGELRLDSPDPQVOKAAQAAVASYNMGSNIYFRDTHIIKAQSLVAGIKYFL 68

Qy 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111

Db 69 TMEMGSTDCRTRVGTGDHVDLT-TCPLAAGAQOEKLCDFEVLVVPWQ 115

RESULT 13

US-08-461-030C-2

Sequence 2, Application US/08461030C

Patent No. 5985601

GENERAL INFORMATION:

APPLICANT: Ni, Jian

APPLICANT: Yu, Guo-Liang

APPLICANT: Gentz, Reiner

APPLICANT: Rosen, Craig A.

TITLE OF INVENTION: Human Cystatin E

NUMBER OF SEQUENCES: 8

CORRESPONDENCE ADDRESS:

ADDRESSEE: Human Genome Sciences, Inc.

STREET: 9410 Key West Ave

CITY: Rockville

STATE: MD

COUNTRY: USA

ZIP: 20850

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent in Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/461.030C

FILING DATE: 05-JUN-1995

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: A. Anders, Brookes

REGISTRATION NUMBER: 36,373

REFERENCE/DOCKET NUMBER: PF202

TELECOMMUNICATION INFORMATION:

TELEPHONE: 301-301-8504

TELEFAX: 301-309-8439

INFORMATION FOR SEQ ID NO: 2:

SEQUENCE CHARACTERISTICS:

LENGTH: 149 amino acids

TYPE: amino acid

TOPOLOGY: linear

MOLECULE TYPE: protein

US-08-461-030C-2

Query Match

Best Local Similarity 20.2%, Score 138.5, DB 2, Length 149,
 Matches 34, Conservative 22, Mismatches 45, Indels 7, Gaps 3;

Qy 10 PTKICVCGPRDIPNPSPELEETLTHITKLNAENNATPYFKIDNVKARVQVAGKKYPI 69
 Db 30 PGRMVGELRLDSPDPQVOKAAQAAVASYNMGSNIYFRDTHIIKAQSLVAGIKYFL 89

OY 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111
DB 90 TWENGSTDCRTRVTGDHVDLT-TCPLAAGAQOEKLRCDFEVLVVPWQ 136

RESULT 14

US-08-744-138-2

Sequence 2, Application US/08744138

Patent No. 6011012

GENERAL INFORMATION:

APPLICANT: Gentz, Reiner L.

APPLICANT: Ni, Jian

APPLICANT: Rosen, Craig A.

APPLICANT: Yu, Guo-Liang

TITLE OF INVENTION: Human Cystatin E

NUMBER OF SEQUENCES: 13

CORRESPONDENCE ADDRESS:

ADDRESSEE: Human Genome Sciences, Inc.

STREET: 9410 Key West Avenue

CITY: Rockville

STATE: Maryland

COUNTRY: USA

ZIP: 20850

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/744,138

FILING DATE:

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: Brookes, A. Anders

REGISTRATION NUMBER: 36,373

REFERENCE/DOCKET NUMBER: PF202P1

TELECOMMUNICATION INFORMATION:

TELEPHONE: 301 309 8504

TELEFAX: 301 309 8512

INFORMATION FOR SEQ ID NO: 2:

SEQUENCE CHARACTERISTICS:

LENGTH: 149 amino acids

TYPE: amino acid

TOPOLOGY: linear

MOLECULE TYPE: protein

US-08-744-138-2

Query Match 20.2%; Score 138.5; DB 3; Length 149;

Best Local Similarity 31.5%; Pred. No. 6.4e-08;

Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

OY 10 PTKICVGCPRDIPNTPNSPELELTHTITKLAENNATFYFKIDNVKARVQVWAGKGYFI 69

DB 30 PQERNVGLRDLSPDDPQVQKAAQAAVASYNMGNSIYYFRDTHIIKAQSQLVAGIKYFL 89

OY 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111

DB 90 TWENGSTDCRTRVTGDHVDLT-TCPLAAGAQOEKLRCDFEVLVVPWQ 136

RESULT 15

US-09-431-480-8

Sequence 8, Application US/09431480

Patent No. 6235708

GENERAL INFORMATION:

APPLICANT: Holloway, James L.

APPLICANT: Feldhaus, Andrew

TITLE OF INVENTION: TESTIS SPECIFIC CYSTATIN-LIKE PROTEIN CYSTATIN T

FILE REFERENCE: 98-72

CURRENT APPLICATION NUMBER: US/09/431,480

CURRENT FILING DATE: 1999-11-01

EARLIER APPLICATION NUMBER: 60/109,217

EARLIER FILING DATE: 1998-11-20

/ EARLIER APPLICATION NUMBER: 60/156,382
/ EARLIER FILING DATE: 1999-09-28
/ NUMBER OF SEQ ID NOS: 22
/ SOFTWARE: FastSeq for Windows Version 3.0
/ SEQ ID NO 8
/ LENGTH: 149
/ TYPE: PRT
/ ORGANISM: Homo sapiens
/ US-09-431-480-8

Query Match

20.2%; Score 138.5; DB 3; Length 149;

Best Local Similarity 31.5%; Pred. No. 6.4e-08;

Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;

OY 10 PTKICVGCPRDIPNTPNSPELELTHTITKLAENNATFYFKIDNVKARVQVWAGKGYFI 69

DB 30 PQERNVGLRDLSPDDPQVQKAAQAAVASYNMGNSIYYFRDTHIIKAQSQLVAGIKYFL 89

OY 70 DFVARETTCSKE-----SNEELTESCETKLGQ--SLDCNAEVVVPWE 111

DB 90 TWENGSTDCRTRVTGDHVDLT-TCPLAAGAQOEKLRCDFEVLVVPWQ 136

Search completed: September 24, 2004, 14:11:37

Job time : 15.732 sec

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: September 24, 2004, 14:05:18 / Search time 36.576 Seconds
(without alignments)
1095.549 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 687

Sequence: 1 GSGKDPVQPTKICVGRD.....VPWKIKIYPTVVAHNECF 127

Scoring table: BLOSUM62

Gapop 10.0 / Gapext 0.5

Searched: 1017041 seqs, 315518202 residues

Total number of hits satisfying chosen parameters: 1017041

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database:

SPTREMBL 251:
1: sp_archaea:
2: sp_bacteria:
3: sp_fungi:
4: sp_human:
5: sp_invertebrate:
6: sp_mammal:
7: sp_mhc:
8: sp_organellar:
9: sp_phage:
10: sp_plant:
11: sp_rodent:
12: sp_virus:
13: sp_vertebrate:
14: sp_unclassified:
15: sp_virus:
16: sp_bacteriap:
17: sp_archaeap:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	383	55.7	140	6 Q7YRP6	Q7YRP6 sus acrofa
2	381	55.5	423	11 P70517	P70517 rattus norv
3	378	55.0	430	11 Q63581	Q63581 rattus norv
4	171.5	25.0	167	11 Q5QWU5	Q5QWU5 mus musculu
5	163.5	23.8	167	4 Q724J8	Q724J8 homo sapien
6	152.5	22.2	462	13 Q72Y91	Q72Y91 xenopus lae
7	152.5	22.2	462	13 Q75YH2	Q75YH2 xenopus lae
8	152.5	22.2	465	13 Q801E5	Q801E5 xenopus lae
9	131.5	19.1	140	11 Q9EPX9	Q9EPX9 mus musculu
10	123.5	18.0	455	13 Q800S8	Q800S8 brachydanio
11	119	17.3	388	11 Q8CB17	Q8CB17 mus musculu
12	117.5	17.1	464	13 Q80125	Q80125 cyprinus ca
13	113.5	16.5	148	5 Q9NH95	Q9NH95 litomossolide
14	113	16.4	140	11 Q80Y72	Q80Y72 mus musculu
15	111	16.2	146	11 Q8K397	Q8K397 mus musculu
16	111	16.2	149	11 Q9D1B1	Q9D1B1 mus musculu

17	108.5	15.8	112	13 Q9SR4	Q9SR4 acipenser s
18	108.5	15.8	112	13 Q9SR3	Q9SR3 acipenser s
19	107	15.6	139	5 Q9TY2	Q9TY2 caenorhabdi
20	106	15.4	300	13 Q80126	Q80126 cyprinus ca
21	105	15.3	109	5 Q9TY65	Q9TY65 onchocerca
22	105	15.3	127	5 P90698	P90698 brugia mala
23	104.5	15.2	149	11 Q8VHC1	Q8VHC1 rattus norv
24	104.5	15.2	161	5 Q16159	Q16159 brugia mala
25	102.5	14.9	127	5 Q9U9A1	Q9U9A1 onchocerca
26	101	14.7	148	11 Q8JMR4	Q8JMR4 mus musculu
27	99	14.4	110	5 Q8SA65	Q8SA65 sandersonia
28	98.5	14.3	107	5 Q8T0Y2	Q8T0Y2 sarcophaga
29	98.5	14.3	125	5 Q25620	Q25620 onchocerca
30	97.5	14.2	134	10 Q41825	Q41825 zea mays (m
31	95.5	13.9	143	5 Q61973	Q61973 caenorhabdi
32	94	13.7	122	5 Q44396	Q44396 haemochus
33	93	13.5	138	4 Q8MXU6	Q8MXU6 homo sapien
34	92	13.4	157	5 Q17108	Q17108 acanthochei
35	90	13.1	139	11 Q8K5A3	Q8K5A3 rattus norv
36	88.5	12.9	92	10 Q9FXN6	Q9FXN6 arabidopsis
37	88.5	12.9	116	10 Q8RXS7	Q8RXS7 arabidopsis
38	88.5	12.9	124	10 Q41906	Q41906 arabidopsis
39	88.5	12.9	125	10 Q22202	Q22202 arabidopsis
40	88.5	12.9	134	10 P93627	P93627 zea mays (m
41	88.5	12.9	134	10 Q41897	Q41897 zea mays (m
42	88	12.8	199	10 Q3270	Q3270 brassica ca
43	87.5	12.7	141	11 Q9DAP1	Q9DAP1 mus musculu
44	86.5	12.6	141	11 Q80ZNS	Q80ZNS mus musculu
45	86.5	12.6	349	6 Q14502	Q14502 cercopithec

ALIGNMENTS

RESULT 1

Q7YRP6 PRELIMINARY; PRT; 140 AA.
AC Q7YRP6
DT 01-OCT-2003 (TRENBLrel. 25, Created)
DT 01-OCT-2003 (TRENBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TRENBLrel. 25, Last annotation update)
DS Low molecular weight kininogen (fragment).
GN KNG.
OS Sus scrofa (Pig).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
OX NCBI_TaxID=9823;
RN [1]
RP SEQUENCE FROM N.A.
RA Vonnahme K.A., Fernando S.C., Ross J.A., Ashworth M.D., Desilva U.,
RA Malayer J.R., Geisert R.D.,
RT "Porcine Endometrial and Conceptus Expression of Kininogens and Plasma
Kallikrein in Cyclic and Pregnant Gilts.",
RL Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY321363; AAP85260.1; -.
FT NON TER 1 140
FT NON TER 1 140
SQ SEQUENCE 140 AA; 15650 MW; 177837836603P777 CRC64;

Query Match 55.7%; Score 383; DB 6; Length 140;
Best Local Similarity 78.9%; Pred. No. 2.8e-30;
Matches 75; Conservative 5; Mismatches 15; Indels 0; Gaps 0;
OY 24 NSPELEETITITIKLNAENNAIPYKIDNKKYQVWVAGKKYFIDFVARETTCSESN 83
DB 1 DSPDLEEPFNHSIAKLNAENNAIPYKIDNKKYQVWVAGKKYFIDFVARETTCSESN 60
OY 84 ERLTSCSTKGLGSLDCNAEVVYVWPKKIYPTV 118
DB 61 ELTSCSTKGLGSLDCNAEVVYVWPKKIYPTV 95

RESULT 2

Anderson K.P., Croyle M.L., Lingrel J.B.;
Primary structure of a gene encoding rat T-kininogen."

RA
RL
Gene 81:119-128 (1989).
EMBL; M29090; AAA42251.1;
DR EMBL; M29083; AAA42251.1; JOINED.
EMBL; M29084; AAA42251.1; JOINED.
DR EMBL; M29091; AAA42251.1; JOINED.
EMBL; M29085; AAA42251.1; JOINED.
DR EMBL; M29086; AAA42251.1; JOINED.
EMBL; M29087; AAA42251.1; JOINED.
DR EMBL; M29088; AAA42251.1; JOINED.
EMBL; M29089; AAA42251.1; JOINED.
PIR; S68034; S68034.
PIR; S68035; S68035.
GO; GO:0004869; Fcycetaeine protease inhibitor activity; IEA.
InterPro; IPR000010; Cystatin.
SMART; PF00031; cystatin; 3.
Pham; SM00043; CY; 3.
PROSITE; PS00287; CYSTATIN; 2.
SEQUENCE 430 AA; 47618 MW; 45508DEF4BDC978C CRC64;

Query Match 55.0%; Score 378; DB 11; Length 430;
Best Local Similarity 62.1%; Pred. No. 3,1e-29;
Matches 72; Conservative 13; Mismatches 31; Indels 0; Gaps 0

OY 3 GKDFVQPPKICVCGRDIPNTSPLEIETLTHTITKLANNATFYFKIDNVKARVOV 62
DB 252 GDOLFELLPNCRGCPREIPVDSPFLKEALGHSHIAQLANQRHHIFYKIDTVAKANSQVV 311

OY 63 AGKYFIDFVARERTCSKSENELTSCSTKKLGSLGLOCNAEVTVVWEKKIVPTV 118
DB 312 AGVIYVFIEAFRTNCQSKXTLDACSTKHGLSCLNCNAVYMRPWNKVPTV 367

RESULT 4
OQOWLS PRELIMINARY; PRT; 167 AA.

ID OQOWLS
AC OQOWLS,
DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)
GN Murine ChAP (CYSTATIN P) (LEUKOCYSTATIN)
OS Mus musculus (Mouse)
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
RN NCBI_TaxId=10090;
RP SEQUENCE FROM N.A.
RR Morita M., Arakawa H., Yoshiuchi N.;
RT "A novel cystatin-like metastasis associated gene,"
RL Submitted (JUN-1998) to the EMBL/GenBank/DDBJ databases.
[2]
SEQUENCE FROM N.A.
RR STRAIN=C57BL/6J; TISSUE=Embryo;
RR MEDLINE=21085660; PubMed=11217851;
RR Kawai Y., Shingnaga A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
ARA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi M., Fukushima S.,
ARA Aizawa K., Okazaki Y., Nishi K., Kiyosawa H., Kondo S., Yamataka I.,
ARA Saito T., Ozaki Y., Colobori T., Bono H., Kasukawa T., Saito R.,
ARA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
ARA Pleischmann W., Gaasterland T., Ghisi C., King B., Kochiwa H.,
ARA Kuehl P., Lewis S., Matsuo Y., Nikaide I., Pesole G., Quackenbush J.,
ARA Schrai L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio T.,
ARA Sakai K., Okido T., Furuno M., Anno H., Baldarelli R., Bash G.,
ARA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
ARA Brownstein M.J., Bult C., Fletcher N., Fujita M., Gariboldi M.,
ARA Gustincich S., Hall D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
ARA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Momabarts P.,
ARA Nordone P., Ring B., Ringenbach C., Rodriguez I., Sakamoto N.,
ARA Sasaki H., Sato K., Schoenbach C., Seva T., Shibata Y., Storch K.-P.,
ARA Suzuki H., Toyooka K., Wang X.H., Waltz C., Whittaker C., Wilming L.,
ARA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohzuki S.,

RA Havaehizaki Y.
RT "Functional annotation of a full-length mouse cDNA collection."
RL Nature 409:685-690(2001).
DR EMBL; AB015224; BAA34940.1; -
DR EMBL; AK004420; BAB23298.1; -
DR HSPF; PU1034; IG96.
DR MGD; MGI:1298217; C97.
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
SQ SEQUENCE 167 AA; 18847 MW; 61F776D8445095FE CRC64;

Query Match 25.0%; Score 171.5; DB 11; Length 167;
Best Local Similarity 35.5%; Pred. No. 3.4e-09;
Matches 39; Conservative 22; Mismatches 42; Indels 7; Gaps 3;

QY 4 KDPVQPTKICVCGPRDIPNPSPELETLTHTITKLAENNAFYFKIDNVKKARVQVVA 63
DB 50 KDIU---SSVKGPPTKTETNPGVLAARSHVEKFNCTNDIFPKSHVSKALVQVVK 106
QY 64 GKXYPIDPVARETTCSKESNEELTESC---TKKLGSLDCNAEYVYVWP 110
DB 107 GLKYLEVKGIRTKCRKTRHQL-DNCDPOTNPALKRTLYCYSEVWVWP 155

RESULT 5
Q724J8 PRELIMINARY; PRT; 167 AA.
AC Q724J8 (TREMBLrel. 25, Created)
DT 01-OCT-2003 (TREMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
DE Cystatin P (leukocystatin).
OS Homo sapiens (human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
[1]
RN SEQUENCE FROM N.A.
RP Kalline N., Chen X., Rolfe A., Halleck A., Hines L., Eisenstein S.,
RA Koundinya M., Raphael J., Moreira D., Kelley T., LaBaer J., Lin Y.,
RA Phelan M., Farmer A.
RL Submitted (AUG-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BT009825; AAP08827.1; -
SQ SEQUENCE 167 AA; 18857 MW; E339025A58D60177 CRC64;

Query Match 23.8%; Score 163.5; DB 4; Length 167;
Best Local Similarity 31.6%; Pred. No. 2.1e-08;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

QY 11 TKICVCGPRDIPNPSPELETLTHTITKLAENNAFYFKIDNVKKARVQVVAQKYPID 70
DB 54 SRVKGPPPTKTNDPGVLAARSHVEKFNCTNDIFPKSHVSKALVQVVKLYML 113
QY 71 FVARETTCSKESNEELTESC---TKKLGSLDCNAEYVYVWPKIYPTVYVHWE 124
DB 114 VEIGRTTCQKQNHRL-DCCDPQNTHTLQTLSCYSEVWVWP-----LQHP 160

RESULT 6
Q72Y91 PRELIMINARY; PRT; 462 AA.
AC Q72Y91 (TREMBLrel. 24, Created)
DT 01-JUN-2003 (TREMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
DE Similar to fetuin B.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipiloidea; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;

RN SEQUENCE FROM N.A.
RP Tissue=Embryo;
RC Klein S., Strauberg R.,
RA Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC043891; AAH43891.1; -
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 2.
DR SMART; SM00043; CY; 2.
SQ SEQUENCE 462 AA; 53185 MW; D7BAD339961739FB CRC64;

Query Match 22.2%; Score 152.5; DB 13; Length 462;
Best Local Similarity 38.8%; Pred. No. 8.4e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;

QY 10 PTKICVCGPRDIPNPSPELETLTHT---ITKLAENNAFYFKIDNVKKARVQVVAQK 65
DB 142 PGVILSTCP-DCPTANEIPTITETADTLIAEYKDSNNTRYFKIDHIERVRSQWVGP 200
QY 66 KYPIDFVARETTCSKESNEELTESC 90
DB 201 SYPIQFTIKETDCMKTOENVLSNC 225

RESULT 7
Q7SYH2 PRELIMINARY; PRT; 462 AA.
AC Q7SYH2 (TREMBLrel. 25, Created)
DT 01-OCT-2003 (TREMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
DE Cystatin domain fetuin-like protein.
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipiloidea; Pipidae;
OC Xenopodinae; Xenopus.
OX NCBI_TaxID=8355;
[1]
RN SEQUENCE FROM N.A.
RP Tissue=Ventral midgut;
RA Costa R.M.B., Mason J., Lee M., Amaya E., Zorn A.M.;
RT "Novel gene expression domains reveal early patterning of the Xenopus
RT endoderm."
RL Submitted (MAR-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY260732; AAP02289.1; -
SQ SEQUENCE 462 AA; 53186 MW; 796P92774CC27721 CRC64;

Query Match 22.2%; Score 152.5; DB 13; Length 462;
Best Local Similarity 38.8%; Pred. No. 8.4e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;

QY 10 PTKICVCGPRDIPNPSPELETLTHT---ITKLAENNAFYFKIDNVKKARVQVVAQK 65
DB 142 PGVILSTCP-DCPTANEIPTITETADTLIAEYKDSNNTRYFKIDHIERVRSQWVGP 200
QY 66 KYPIDFVARETTCSKESNEELTESC 90
DB 201 SYPIQFTIKETDCMKTOENVLSNC 225

RESULT 8
Q801E5 PRELIMINARY; PRT; 465 AA.
AC Q801E5 (TREMBLrel. 24, Created)
DT 01-JUN-2003 (TREMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
DE Hypothetical histidine-rich protein (fragment).
OS Xenopus laevis (African clawed frog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Anura; Mesobatrachia; Pipiloidea; Pipidae;
OC Xenopodinae; Xenopus.

```

OX NCBI_TaxID=8355;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22480013; PubMed=12591597;
RA Chen Y., Jurgens K., Hollemann T., Clausen M., Ramadori G.,
RA Pieler T.,
RT "Cell-autonomous and signal-dependent expression of liver and
RT intestine marker genes in pluripotent precursor cells from Xenopus
RT embryos.",
RL Mech. Dev. 120:277-288(2003).
DR EMBL: AY188284; AAC01610.1; -.
DR GO: GO:0004869; P:cysteine protease inhibitor activity; IEA.
DR InterPro: IPR000010; Cystatin.
DR Pfam: PF00031; Cystatin; 2.
DR SMART: SM00043; Cy; 2.
KW Hypothetical protein.
SQ
SEQUENCE 465 AA; 53528 MW; 0B403AB4P78BBFD4 CRC64;

Query Match 22.24; Score 152.5; DB 13; Length 465;
Best Local Similarity 38.84; Pred. No. 8.5e-07;
Matches 33; Conservative 13; Mismatches 34; Indels 5; Gaps 2;

OY 10 PTKLCVCPDIPDTPNSPELETLTHT---ITKLAENNAATFFKIDNVKARVOVWAGK 65
DB 145 PGVILSTCP-DCTANEELTPITETAEFLIAEYKDSNTRIFKIDHIERVRSQWVGP 203

OY 66 KYPIDFVARETTCKESNEELTESC 90
DB 204 SYFIQPTIKSTOCTCKTOENVLNSC 228

RESULT 9
ID Q9EPX9 PRELIMINARY; PRT; 140 AA.
AC Q9EPX9
DT 01-MAR-2001 (TrEMBLrel. 16, Created)
DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DR Cystatin C.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21010502; PubMed=11143350;
RA Taupin P.J., Ray J., Fischer W.H., Suhr S.T., Hakansson K., Grubb A.,
RA Gage P.H.;
RT "FGF-2-Responsive neural stem cell proliferation requires CCG, a novel
RT autocrine/paracrine cofactor.",
RL Neuron 28:385-397(2000).
DR EMBL: AF111741; AAC40283.1; -.
DR HSP: P01034; ICG6.
DR GO: GO:0004869; P:cysteine protease inhibitor activity; IEA.
DR InterPro: IPR000010; Cystatin.
DR Pfam: PF00031; Cystatin; 1.
DR SMART: SM00043; Cy; 1.
DR PROSITE: PS00287; CYSTATIN; 1.
CHAIN 21 140 CYSTATIN C.
FT VARIANTS 16 16 A -> G.
FT VARIANT 84 84 L -> F.
SQ
SEQUENCE 140 AA; 15517 MW; 3A563406DD58D785 CRC64;

Query Match 19.14; Score 131.5; DB 11; Length 140;
Best Local Similarity 27.84; Pred. No. 2.6e-05;
Matches 32; Conservative 26; Mismatches 48; Indels 9; Gaps 4;

OY 15 VGCPRDIPDTPNSPELETLTHTITKLAENNAATFFKIDNVKARVOVWAGKYPIDFVAR 74
DB 30 LGAPPEADANECCVRALDFAVSEYNKGSNDAYHSRAIQVVRARQQLVAGVNYFLDVMG 89

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OY 75 ETTCKESNEELTESC---ETKLGQSLDCNAEVVVVWPEKKIYPTVTWNHCE 126
DB 90 RITCTK-SQTNLTD-CPPHDQPHLMKALCSFOIYSPWK----GTHSLTNPSCK 138

RESULT 10
ID Q800S8 PRELIMINARY; PRT; 455 AA.
AC Q800S8
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DR Fetuin-A.
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Osteichthyes; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
RA Jia F.;
RT "Danio rerio fetuin-A.",
RL Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL: AY217758; AAC61483.1; -.
DR GO: GO:0005874; C:microtubule; IEA.
DR GO: GO:0004869; P:cysteine protease inhibitor activity; IEA.
DR GO: GO:0005198; P:structural molecule activity; IEA.
DR GO: GO:0007018; P:microtubule-based movement; IEA.
DR InterPro: IPR002453; Beta tubulin.
DR InterPro: IPR000010; Cystatin.
DR Pfam: PF00031; Cystatin; 1.
DR SMART: SM00043; Cy; 1.
DR PROSITE: PS00228; TUBULIN B AUTOREG; 1.
SQ
SEQUENCE 455 AA; 50627 MW; D822872926BAJACB CRC64;

Query Match 18.04; Score 123.5; DB 13; Length 455;
Best Local Similarity 26.74; Pred. No. 0.00063;
Matches 32; Conservative 23; Mismatches 46; Indels 19; Gaps 4;

OY 2 SGHDFVQPTKICVGCPRDIPDTPNSPELETLTHTITKLAENNAATFFKIDNVKARVQ- 60
DB 134 SHEDLV---KKCPDCGGLFLHPEKALSVNAALAKFKSNHKSYPKLMVGRISQW 189

OY 61 VVAGKYPIDFVARETTCKESNEELTESC-----CETKLG-OSLDCNAEVY 106
DB 190 MPMGOSYPTQFAIMETNCTKKDAPQNPPEACKALCGDQATYGFCKSKVSGSEPEVECIY 249

RESULT 11
ID Q8CB17 PRELIMINARY; PRT; 388 AA.
AC Q8CB17
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DR Fetuin beta.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Vagina;
RX MEDLINE=22354883; PubMed=12466851;
RA The FANTOM Consortium,
RA the RIKEN Genome Exploration Research Group Phase I & II Team;
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs.",
RL Nature 420:563-573(2002).
DR EMBL: AK037043; BAC29682.1; -.
DR MGI:1890221; Fetub.

```


RP SEQUENCE FROM N.A.
RC TISSUE=estricle;
RA Strausberg R.;
RL Submitted (MAR-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC048646; AAH48646.1; -;
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR DR InterPro; IPR003243; Cystatin_C/M.
DR Pfam; PF00031; Cystatin; 1.
DR ProDom; PD001231; Cystatin_C/M; 1.
DR SMART; SM00043; CY; 1.
SQ SEQUENCE 140 AA; 16199 MW; 32633B99C4697DA0 CRC64;

Query Match 16.4%; Score 113; DB 11; Length 140;
Best Local Similarity 29.4%; Pred. No. 0.0018;
Matches 25; Conservative 18; Mismatches 38; Indels 4; Gaps 2;

QY 28 LEETLTIITKLNAENNAATPKIDNVKARVQVAGKKYFIDFVARETTCSKESNELT 87
DB 44 INSTLHPFIRSYNASNDLYIQVQKLIQGMQLTGVYLVTVKIGRTCKK--NETKK 101
QY 88 ESC--ETKQLGQSLDCNAEVVVPW 110
DB 102 ASCPLQSSKLKSLICKSLIYSVPW 136

RESULT 15
QY 08K397 PRELIMINARY; PRT; 146 AA.
AC 08K397,
DT 01-OCT-2002 (TrEMBLrel. 22, Created)
DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
DE 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE RIKEN CDNA 110017811 gene (fragment).
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Mammary gland;
RA Strausberg R.;
RL Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC027680; AAH27680.1; -;
DR GO; GO:0004869; F:cysteine protease inhibitor activity; IEA.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 1.
DR SMART; SM00043; CY; 1.
PT NON TER 1
SQ SEQUENCE 146 AA; 16380 MW; 9D77BB9A6063A5C4 CRC64;

Query Match 16.2%; Score 111; DB 11; Length 146;
Best Local Similarity 30.5%; Pred. No. 0.003;
Matches 29; Conservative 18; Mismatches 42; Indels 6; Gaps 4;

QY 22 PTHSPLESELTHTITKLNAENNAATPKIDNVKARVQVAGKKYFIDFVARETTCSKE 81
DB 40 PTD-PRVQAQAQAAVASTNGSDSLYPRDTKVIDAKYLVAGIKYLLDIESTECKT 98
QY 82 --SNEEL-TESCETKQLQ--SLDCNAEVVVPWE 111
DB 99 RVSGEHMDLTTCPLAAGGQOEKRCNPELLEVPWK 133

Search completed: September 24, 2004, 14:10:18
Job time : 37.576 secs

GenCore version 5.1.6
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OK protein - protein search, using sw model

Run on: September 24, 2004, 14:04:32 ; Search time 8.636 seconds
(without alignment)
765.738 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 687

Sequence: 1 GSGKDFVGPPTKICVGCPRD.....VPWEKKIYPTVTNHRCEP 127

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 141681 seqs, 52070155 residues

Total number of hits satisfying chosen parameters: 141681

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_42.4

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Query Length	ID	Description
1	618	90.0	644	1 KNG_HUMAN	P01042 homo sapien
2	440	64.0	436	1 KXLI_BOVIN	P01046 bos taurus
3	440	64.0	621	1 KXHI_BOVIN	P01044 bos taurus
4	413	60.1	434	1 KXLI_BOVIN	P01047 bos taurus
5	413	60.1	619	1 KXH2_BOVIN	P01047 bos taurus
6	413	60.1	661	1 KNG_MOUSE	O08677 mus musculus
7	410	59.7	639	1 KNG_RAT	P08934 rattus norv
8	388	56.5	430	1 KNT2_RAT	P08932 rattus norv
9	380	55.3	430	1 KNT1_RAT	P01048 rattus norv
10	171.5	25.0	144	1 CYTF_MOUSE	O89098 mus musculus
11	163.5	23.8	145	1 CYTF_HUMAN	O76096 homo sapien
12	138.5	20.2	146	1 CYTC_MACMU	O19092 macaca mlla
13	138.5	20.2	149	1 CYTM_HUMAN	O15928 homo sapien
14	137.5	20.0	148	1 CYTC_BOVIN	P01035 bos taurus
15	135	19.7	378	1 FETB_RAT	O9QX79 rattus norv
16	132.5	19.3	146	1 CYTC_HUMAN	P01034 homo sapien
17	132.5	19.3	146	1 CYTC_SALISC	O19093 salmisi sci
18	131.5	19.1	382	1 FETB_HUMAN	O9QUM5 homo sapien
19	130	18.9	127	1 CYTC_RAT	P14841 rattus norv
20	129.5	18.9	140	1 CYTC_MOUSE	P21460 mus musculus
21	128	18.6	111	1 CYT_BITAR	P08935 bitis ariet
22	124.5	18.1	141	1 CYTT_HUMAN	O97228 homo sapien
23	124.5	18.1	148	1 CYTT_RABIT	O97862 oryctolagus
24	122.5	17.8	116	1 CYT_COTUA	P81061 coturnix co
25	119	17.3	388	1 FETB_MOUSE	O9QXCI mus musculus
26	118.5	17.2	139	1 CYT_CHICK	P01038 gallus gall
27	113	16.4	141	1 CYTS_RAT	P19313 rattus norv
28	109.5	15.9	141	1 CYTN_HUMAN	P01037 homo sapien
29	108.5	15.8	141	1 CYTS_HUMAN	P01036 homo sapien
30	107	15.6	130	1 CYT_ONCKE	O98967 oncorhynch
31	105.5	15.4	162	1 CYTX_ONCVO	P22085 onchocerca
32	105	15.3	130	1 CYT_ONCKY	O91195 oncorhynch
33	104	15.1	129	1 CYT_CYPCA	P35481 cyprinus ca

RESULT 1

KNG_HUMAN STANDARD; PRT; 644 AA.
AC P01042, P01043;
DT 21-JUL-1986 (Rel. 01, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Kinogen precursor (Alpha-2-thiol proteinase inhibitor) [Contains:
DE Bradykinin].
GN KNG.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORMS HMW AND LMW).
RC TISSUE=Liver;
RX MEDLINE=85234582; PubMed=2989293;
RA Takagaki Y., Kitamura N., Nakanishi S.;
RT "Cloning and sequence analysis of cDNAs for human high molecular
RT weight and low molecular weight prekininogens. Primary structures of
RT two human prekininogens.";
RL J. Biol. Chem. 260:8601-8609 (1985).
RN [2]
RP GENE STRUCTURE.
RX MEDLINE=85234583; PubMed=2989294;
RA Kitamura N., Kitagawa H., Fukushima D., Takagaki Y., Miyata T.,
RA Nakanishi S.;
RT "Structural organization of the human kininogen gene and a model for
RT its evolution.";
RL J. Biol. Chem. 260:8610-8617 (1985).
RN [3]
RP SEQUENCE OF 1-401 FROM N.A.
RX MEDLINE=85122621; PubMed=6441591;
RA Okubo I., Kurachi K., Takasawa T., Shiohawa H., Sasaki M.;
RT "Isolation of a human cDNA for alpha 2-thiol proteinase inhibitor and
RT its identity with low molecular weight kininogen.";
RL Biochemistry 23:5691-5697 (1984).
RN [4]
RP SEQUENCE OF 379-644.
RX MEDLINE=86030270; PubMed=4054110;
RA Lottspeich F., Kellermann J., Henschen A., Foerster B.,
RA Mueller-Eberhard W.;
RT "The amino acid sequence of the light chain of human high-molecular-
RT mass kininogen.";
RL Eur. J. Biochem. 152:307-314 (1985).
RN [5]
RP SEQUENCE OF 381-389.
RX MEDLINE=90255622; PubMed=4952632;
RA Pierce J.V.;
RT "Structural features of plasma kinins and kininogens.";
RL Fed. Proc. 27:52-57 (1968).
RN [6]
RP DISULFIDE BONDS.
RA Sueyoshi T., Miyata T., Kato H., Iwanaga S.;
RT "Disulfide bonds in bovine HMW kininogens.";

ALIGNMENTS

34 102 14.8 122 1 CYTA_SARPE
35 101 14.7 139 1 CS11_MOUSE
36 95 13.8 165 1 CSTL_HUMAN
37 94.5 13.8 142 1 CST8_MOUSE
38 94 13.8 345 1 A2HS_MOUSE
39 94 13.7 142 1 CST8_HUMAN
40 93 13.5 99 1 CYT_NAJAT
41 93 13.5 137 1 CS11_HUMAN
42 93 13.5 142 1 CYTD_HUMAN
43 91.5 13.3 135 1 CYTD_MAIZE
44 91.5 13.3 352 1 A2HS_RAT
45 88 12.8 367 1 A2HS_HUMAN

P31727 sarcophaga
Q9d269 mus musculus
Q9h114 homo sapien
P32766 mus musculus
P39699 mus musculus
O60676 homo sapien
P81714 naia atra
Q9h112 homo sapien
P38335 homo sapien
P31786 sca maye (m
P24090 rattus norv
P02765 homo sapien

RL Seikagaku 56:808-808(1984).
 RN [7]
 RP CARBOHYDRATE-LINKAGE SITE ASN-294.
 RX MEDLINE=22660472; PubMed=12754519;
 RA Zhang H., Li X.-J., Martin D.B., Abersold R.,
 RT "Identification and quantification of N-linked glycoproteins using
 RT hydrazide chemistry, stable isotope labeling and mass spectrometry.";
 RL Nat. Biotechnol. 21:660-666(2003).
 CC -1- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
 CC HMW-kininogen plays an important role in blood coagulation by
 CC helping to position optimally prekallikrein and factor XI next to
 CC factor XII; (3) HMW-kininogen inhibits the thrombin and plasmin-
 CC induced aggregation of thrombocytes; (4) the active peptide
 CC bradykinin that is released from HMW-kininogen shows a variety of
 CC physiological effects: (4A) influence in smooth muscle
 CC contraction, (4B) induction of hypotension, (4C) natriuresis and
 CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
 CC mediator of inflammation and causes (4E1) increase in vascular
 CC permeability, (4E2) stimulation of nociceptors (4E3) release of
 CC other mediators of inflammation (e.g. prostaglandins). (4F) it has
 CC a cardioprotective effect (directly via bradykinin action,
 CC indirectly via endothelium-derived relaxing factor action); (5)
 CC HMW-kininogen inhibits the aggregation of thrombocytes; (6) LMW-
 CC kininogen is in contrast to HMW-kininogen not involved in blood
 CC clotting.
 CC -1- SUBCELLULAR LOCATION: Secreted.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=2;
 CC Name=HMW;
 CC IsoId=P01042-1; Sequence=Displayed;
 CC Name=LMW;
 CC IsoId=P01042-2; Sequence=VSP_001261; VSP_001262;
 CC -1- TISSUE SPECIFICITY: Plasma.
 CC -1- PTM: Bradykinin is released from kininogen by plasma kallikrein.
 CC -1- SIMILARITY: Contains 3 cystatin-like domains.
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
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 CC or send an email to license@sib-sib.ch).
 CC -----
 CC EMBL: K02566; AAB35497.1; -
 CC EMBL: M11437; AAB59550.1; -
 CC EMBL: M11438; AAB59550.1; JOINED.
 CC EMBL: M11521; AAB59550.1; JOINED.
 CC EMBL: M11522; AAB59550.1; JOINED.
 CC EMBL: M11523; AAB59550.1; JOINED.
 CC EMBL: M11524; AAB59550.1; JOINED.
 CC EMBL: M11525; AAB59550.1; JOINED.
 CC EMBL: M11526; AAB59550.1; JOINED.
 CC EMBL: M11527; AAB59550.1; JOINED.
 CC EMBL: M11528; AAB59550.1; JOINED.
 CC EMBL: M11437; AAB59551.1; -
 CC EMBL: M11438; AAB59551.1; JOINED.
 CC EMBL: M11521; AAB59551.1; JOINED.
 CC EMBL: M11522; AAB59551.1; JOINED.
 CC EMBL: M11523; AAB59551.1; JOINED.
 CC EMBL: M11524; AAB59551.1; JOINED.
 CC EMBL: M11525; AAB59551.1; JOINED.
 CC EMBL: M11526; AAB59551.1; JOINED.
 CC EMBL: M11527; AAB59551.1; JOINED.
 CC EMBL: M11528; AAB59551.1; JOINED.
 CC PIR: A01279; KGHUHI.
 CC FIC: A01280; KGHUHI.
 CC SWISS-2DPAGE: P01042; HUMAN.
 CC GeneW: HGNC:6383; KNG.
 CC MIN: 228960; -
 CC GO: GO:0007596; P: blood coagulation; NAS.
 CC GO: GO:0030146; P: diuresis; NAS.
 CC GO: GO:0006954; P: inflammatory response; NAS.

DR GO: GO:0030147; P: natriuresis; NAS.
 DR GO: GO:0006939; P: smooth muscle contraction; NAS.
 DR InterPro: IPR000010; Cystatin.
 DR InterPro: IPR002395; Kininogen.
 DR Pfam: PF00031; cystatin_3.
 DR PRINTS: PR00334; KININOGEN.
 DR SMART: SM00043; CY_3.
 DR PROSITE: PS00287; CYSTATIN_2.
 KW Glycoprotein; plasma; repeat; Thiol protease inhibitor; Vasodilator;
 KW Bradykinin; Blood coagulation; Inflammatory response; Signal;
 KW Alternative splicing; Pyrrolidone carboxylic acid.
 FT SIGNAL 1 18
 FT CHAIN 19 644 KININOGEN.
 FT CHAIN 19 380 KININOGEN HEAVY CHAIN.
 FT PEPTIDE 381 389 BRADYKININ.
 FT CHAIN 390 644 KININOGEN LIGHT CHAIN.
 FT DOMAIN 19 136 CYSTATIN-LIKE 1.
 FT DOMAIN 137 258 CYSTATIN-LIKE 2.
 FT DOMAIN 259 380 CYSTATIN-LIKE 3.
 FT DOMAIN 420 510 HIS-RICH
 FT REPEAT 420 449 (ASSOCIATED WITH CLOTTING ACTIVITY).
 FT REPEAT 450 479
 FT REPEAT 480 510
 FT MOD_RES 19 19
 FT DISULFID 28 614
 FT DISULFID 83 94
 FT DISULFID 107 126
 FT DISULFID 142 145
 FT DISULFID 206 218
 FT DISULFID 229 248
 FT DISULFID 264 267
 FT DISULFID 328 340
 FT DISULFID 351 370
 FT CARBOHYD 48 48
 FT CARBOHYD 169 169
 FT CARBOHYD 205 205
 FT CARBOHYD 294 294
 FT CARBOHYD 401 401
 FT CARBOHYD 533 533
 FT CARBOHYD 542 542
 FT CARBOHYD 546 546
 FT CARBOHYD 557 557
 FT CARBOHYD 571 571
 FT CARBOHYD 577 577
 FT CARBOHYD 593 593
 FT CARBOHYD 628 628
 FT VARSPLIC 402 427
 FT VARSPLIC 428 644
 FT CONFLICT 593 593
 FT SEQUENCE 644 AA; 71945 MW; 313284CBAP8PB87E CRC64;
 Query Match 90.0%; Score 618; DB 1; Length 644;
 Best Local Similarity 100.0%; Pred. No. 3.4e-51;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GDFVPPPTKICVCPEDIPNPSPELETTHITIKNAENNTAFYKIDNVKARVQV 62
 DB 253 GDFVPPPTKICVCPEDIPNPSPELETTHITIKNAENNTAFYKIDNVKARVQV 312
 QY 63 ACKKTFIDFVARETTCSKESNEELTSCETKLGQSLDCNAEYVVPWEKKIYPTV 118
 DB 313 ACKKTFIDFVARETTCSKESNEELTSCETKLGQSLDCNAEYVVPWEKKIYPTV 368
 RESULT 2
 ID KN11 BOVIN STANDARD; PRT; 436 AA.
 AC P01046;
 DT 21-JUL-1986 (Rel. 01, Created)

21-JUL-1986 (Rel. 01, Last sequence update)
 15-MAR-2004 (Rel. 43, Last annotation update)
 Kininogen, LMW I precursor (Thiol proteinase inhibitor) (Contains: Bradykinin).
 Bos taurus (Bovine).
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovinae; Bos.
 NCBI_TaxID=9913;
 [1]
 RN SEQUENCE FROM N.A.
 RP MEDLINE=83117859; PubMed=6572010;
 RA Nawa H., Kitamura N., Hirose T., Asai M., Inayama S., Nakanishi S.;
 RT "Primary structures of bovine liver low molecular weight kininogen
 precursors and their two mRNAs";
 RL Proc. Natl. Acad. Sci. U.S.A. 80:90-94 (1983).
 RN [2]
 RP SEQUENCE OF 19-378.
 RX MEDLINE=87137530; PubMed=3546295;
 RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
 Miyata T., Iwanaga S.;
 RT "Bovine high molecular weight kininogen. The amino acid sequence,
 positions of carbohydrate chains and disulfide bridges in the heavy
 chain portion.";
 RL J. Biol. Chem. 262:2768-2779 (1987).
 CC -1- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
 CC LMW-kininogen inhibits the aggregation of thrombocytes; (3) the
 CC active peptide kallidin that is released from LMW-kininogen shows
 CC a variety of physiological effects: (3A) influence in smooth
 CC muscle contraction, (3B) induction of hypotension, (3C)
 CC natriuresis and diuresis (kidney).
 CC -1- SUBCELLULAR LOCATION: Extracellular.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Event=alternative splicing; Named isoforms=2;
 CC Name=LMW I;
 CC IsoId=IP01046-1; Sequence=Displayed;
 CC Name=HMW I;
 CC IsoId=IP01044-1; Sequence=External;
 CC -1- TISSUE SPECIFICITY: Plasma.
 CC -1- PTM: Bradykinin is released from kininogen by plasma kallikrein.
 CC -1- MISCELLANEOUS: LMW-kininogen is in contrast to HMW-kininogen not
 CC involved in blood clotting.
 CC -1- SIMILARITY: Contains 3 cystatin-like domains.
 CC
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 CC
 CC EMBL; V00426; CAA23709.1; -.
 DR PIR; A01283; KGBOL1.
 DR InterPro; IPR000010; Cystatin.
 DR Pfam; PF00031; cystatin; 3.
 DR SMART; SM00043; CV; 3.
 DR PROSITE; PS00287; CYSTATIN; 2.
 KW Glycoprotein; plasma; Repeat; Vasodilator; Alternative splicing;
 KW Thiol protease inhibitor; Bradykinin; Signal;
 KW Pyroglutamate carboxylic acid.
 FT SIGNAL 1 18
 FT CHAIN 19 436 KININOGEN, LMW I.
 FT CHAIN 19 378 HEAVY CHAIN.
 FT PEPTIDE 380 388 BRADYKININ.
 FT CHAIN 389 436 LIGHT CHAIN.
 FT DOMAIN 19 135 CYSTATIN-LIKE 1.
 FT DOMAIN 136 257 CYSTATIN-LIKE 2.
 FT DOMAIN 258 378 CYSTATIN-LIKE 3.
 FT MOD RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
 FT MOD RES 87 87 N-LINKED (GLCNAC. . .).
 FT CARBOHYD 87 87 O-LINKED (PARTIAL).
 FT CARBOHYD 136 136 O-LINKED (PARTIAL).
 FT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).

FT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
 FT CARBOHYD 204 204 N-LINKED (GLCNAC. . .).
 FT DISULFID 27 406 INTERCHAIN.
 FT DISULFID 82 93
 FT DISULFID 106 125
 FT DISULFID 141 144
 FT DISULFID 205 217
 FT DISULFID 228 247
 FT DISULFID 263 266
 FT DISULFID 327 339
 FT DISULFID 350 369
 FT CONFLICT 295 295
 FT SEQUENCE 436 AA; 48427 MW; P01F78B6814BCE6C CXC64;
 SQ
 Query Match 54.0%; Score 440; DB 1; Length 436;
 Best Local Similarity 70.4%; Pred. No. 1.8e-34;
 Matches 81; Conservative 14; Mismatches 20; Indels 0; Gaps 0;
 QY 4 KDPVQPTKICVCGPRDIPFNSEPELEETITHTITKLNANNATYPIKIDNVKARVQVVA 63
 DB 253 KDPVQPTKICVCGPRDIPFNSEPELEETITHTITKLNANNATYPIKIDNVKARVQVVA 312
 QY 64 GKYPIDPVARETTCSEKNEELTSCETKKGSLGSLDCNARFVYVVPWEKKIYPTV 118
 DB 313 GKYSIVFIARETTCSKSGNEELTKSCINITHQILKCDANVYVVPWEKKIYPTV 367
 RESULT 3
 ID KWHI BOVIN STANDARD; PRT; 621 AA.
 AC P01044;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 21-JUL-1986 (Rel. 01, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Kininogen, HMW I precursor (Thiol proteinase inhibitor) (Contains:
 DE Bradykinin).
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 OK NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=84014106; PubMed=6571699;
 RA Kitamura N., Takagaki Y., Furuto S., Tanaka T., Nawa H., Nakanishi S.;
 RT "A single gene for bovine high molecular weight and low molecular
 RT weight kininogens";
 RL Nature 305:545-549 (1983).
 RN [2]
 RP SEQUENCE OF 19-378.
 RX MEDLINE=87137530; PubMed=3546295;
 RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
 Miyata T., Iwanaga S.;
 RT "Bovine high molecular weight kininogen. The amino acid sequence,
 RT positions of carbohydrate chains and disulfide bridges in the heavy
 RT chain portion.";
 RL J. Biol. Chem. 262:2768-2779 (1987).
 RN [3]
 RP SEQUENCE OF 378-393.
 RX MEDLINE=70180420; PubMed=4986212;
 RA Kato H., Nagasawa S., Suzuki T.;
 RT "Studies on the structure of bovine kininogen: cleavages of disulfide
 RT bonds and of methionyl bonds in kininogen-II";
 RL J. Biochem. 67:313-323 (1970).
 RN [4]
 RP SEQUENCE OF 458-498.
 RX MEDLINE=75170265; PubMed=1169237;
 RA Han Y.N., Komiya M., Iwanaga S., Suzuki T.;
 RT "Studies on the primary structure of bovine high-molecular-weight
 RT kininogen. Amino acid sequence of a fragment ('histidine-rich
 RT peptide') released by plasma kallikrein";
 RL J. Biochem. 77:55-68 (1975).
 CC -1- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)

Thiol protease inhibitor; Bradykinin; Signal;
 Pyroglutamate carboxylic acid.
 KW SIGNAL 1 18
 FT CHAIN 19 434 KININOGEN, LMW II.
 FT PEPTIDE 378 396 HEAVY CHAIN.
 FT CHAIN 387 434 BRADYKININ.
 FT CHAIN 387 434 LIGHT CHAIN.
 FT CHAIN 387 434 CYPSTATIN-LIKE 1.
 FT CHAIN 387 434 CYPSTATIN-LIKE 2.
 FT CHAIN 387 434 CYPSTATIN-LIKE 3.
 FT CHAIN 387 434 PYROGLUTAMATE CARBOXYLIC ACID.
 FT CHAIN 387 434 N-LINKED (GLCNAC).
 FT CHAIN 387 434 O-LINKED (GLCNAC).
 FT CHAIN 387 434 N-LINKED (GLCNAC).
 FT CHAIN 387 434 N-LINKED (GLCNAC).
 FT CHAIN 387 434 INTERCHAIN.
 SQ SEQUENCE 434 AA; 48148 MW; 73A7079DE3E03430 CRC64;
 Query Match 60.1%; Score 413; DB 1; Length 434;
 Best Local Similarity 79.2%; Pred. No. 6.6e-32;
 Matches 79; Conservative 14; Mismatches 22; Indels 2; Gaps 1;
 QY 3 GKDVPPTKICVGPDPIDPTNPSPELTHTTKLNARNATVPKIDNVKARQVQV 62
 DB 252 GEPFL--PPWCVGCPKPIVDSPDLNLSAKLNASHDGTFFKIDTVKATQVQV 309
 QY 63 AGKYPIDFVARETTCSEKSELTETSCETKGLQSLDCAEYVYVVEKIKYPTV 118
 DB 310 GGLKYSILVPIARETTCSEKSELTETSCETKGLQSLDCAEYVYVVEKIKYPTV 365
 RESULT 5
 KMH2 BOVIN STANDARD; PRT; 619 AA.
 ID KMH2 BOVIN STANDARD; PRT; 619 AA.
 AC P01045;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 21-JUL-1986 (Rel. 01, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Kininogen, HMW II precursor (thiol proteinase inhibitor) (Contains:
 Bradykinin).
 OS Bos taurus (Bovinae).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Butheraia; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 OX NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=84014106; PubMed=6571699;
 RA Kitamura N., Takagaki Y., Furuta S., Tanaka T., Nawa H., Nakanishi S.,
 "A single gene for bovine high molecular weight and low molecular
 weight kininogens",
 FT weight kininogens",
 RL Nature 305:545-549 (1983).
 RN [2]
 RP SEQUENCE OF 19-376.
 RX MEDLINE=87137530; PubMed=3546295;
 RA Sueyoshi T., Miyata T., Hashimoto N., Kato H., Hayashida H.,
 Miyata T., Iwanaga S.,
 "Bovine high molecular weight kininogen. The amino acid sequence,
 positions of carbohydrate chains and disulfide bridges in the heavy
 chain portion.",
 FT chain portion.",
 RL J. Biol. Chem. 262:2768-2779 (1987).
 RN [3]
 RP SEQUENCE OF 376-391.

MEDLINE=70180420; PubMed=4986212;
 Kato H., Nagaawa S., Suzuki T.,
 "Studies on the structure of bovine kininogen: cleavages of disulfide
 bonds and of methionyl bonds in kininogen-II.",
 J. Biochem. 67:313-323 (1970).
 [4]
 SEQUENCE OF 387-455.
 MEDLINE=76260155; PubMed=956151;
 Han Y.N., Kato H., Iwanaga S., Suzuki T.,
 "Primary structure of bovine plasma high-molecular-weight kininogen.
 The amino acid sequence of a glycopeptide portion (fragment 1)
 following the C-terminus of the bradykinin moiety.",
 J. Biochem. 79:1201-1222 (1976).
 [5]
 SEQUENCE OF 456-496.
 MEDLINE=75170365; PubMed=1169237;
 Han Y.N., Komiya M., Iwanaga S., Suzuki T.,
 "Studies on the primary structure of bovine high-molecular-weight
 kininogen. Amino acid sequence of a fragment ('histidine-rich
 peptide') released by plasma kallikrein.",
 J. Biochem. 77:55-68 (1975).
 -1- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
 HMW-kininogen plays an important role in blood coagulation by
 helping to position optimally prekallikrein and factor XI next to
 factor XII; (3) HMW-kininogen inhibits the thrombin- and plasmin-
 induced aggregation of thrombocytes; (4) the active peptide
 bradykinin that is released from HMW-kininogen shows a variety of
 physiological effects: (4A) influence in smooth muscle
 contraction, (4B) induction of hypotension, (4C) natriuresis and
 diuresis, (4D) decrease in blood glucose level, (4E) it is a
 mediator of inflammation and causes (4E1) increase in vascular
 permeability, (4E2) stimulation of nociceptors (4E3) release of
 other mediators of inflammation (e.g. prostaglandins), (4F) it has
 a cardioprotective effect (directly via bradykinin action,
 indirectly via endothelium-derived relaxing factor action).
 -1- SUBCELLULAR LOCATION: Extracellular.
 -1- ALTERNATIVE PRODUCTS:
 Name=HMW II;
 IsoId=P01045-1; Sequence=Displayed;
 Name=LMW II;
 IsoId=P01047-1; Sequence=External;
 -1- TISSUE SPECIFICITY: Plasma.
 -1- PTM: Bradykinin is released from kininogen by plasma kallikrein.
 -1- SIMILARITY: Contains 3 cystatin-like domains.
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 EMBL, V01492; CAA24736.1;
 PIR, A01282; K0B0H2.
 HSP, P01038; I490.
 InterPro: IPR000010; Cystatin.
 InterPro: IPR002395; Kininogen.
 Pfam: PF00031; cystatin; 3.
 PRINTS: PR00334; KININOGEN.
 SMART: SM00043; CV, 3.
 PROSITE: PS00387; Cystatin; 2
 Glycoprotein; Plasma Repeat; vasodilator; Alternative splicing;
 Thiol protease inhibitor; Bradykinin; blood coagulation; Signal;
 Inflammatory response; Pyroglutamate carboxylic acid.
 FT SIGNAL 1 18
 FT CHAIN 19 619 KININOGEN, HMW II.
 FT CHAIN 19 376 HEAVY CHAIN.
 FT PEPTIDE 378 386 BRADYKININ.
 FT CHAIN 387 619 LIGHT CHAIN.
 FT CHAIN 387 619 CYPSTATIN-LIKE 1.
 FT CHAIN 387 619 CYPSTATIN-LIKE 2.
 FT CHAIN 387 619 CYPSTATIN-LIKE 3.

PT DOMAIN 257 376 CYPSTATIN-LIKE 3.
 PT MOD_RES 19 19 PYRROLIDONE CARBOXYLIC ACID.
 PT CARBOHYD 87 87 N-LINKED (GLCNAC. . .)
 PT CARBOHYD 136 136 O-LINKED (PARTIAL).
 PT CARBOHYD 168 168 N-LINKED (GLCNAC. . .) (OR 169).
 PT CARBOHYD 197 197 N-LINKED (GLCNAC. . .) (PARTIAL).
 PT CARBOHYD 204 204 N-LINKED (GLCNAC. . .)
 PT CARBOHYD 280 280 N-LINKED (GLCNAC. . .)
 PT CARBOHYD 400 400 O-LINKED.
 PT DISULFID 27 589 INTERCHAIN.
 PT DISULFID 82 93
 PT DISULFID 106 125
 PT DISULFID 141 144
 PT DISULFID 205 217
 PT DISULFID 228 247
 PT DISULFID 261 264
 PT DISULFID 325 337
 PT DISULFID 348 367
 PT VARIANT 398 398
 PT VARIANT 401 401
 PT VARIANT 454 454
 PT SEQUENCE 619 AA, 68710 MW, P04320AB80E0E0DA CRC64;
 Query Match 60.1%; Score 413; DB 1; Length 619;
 Best Local Similarity 67.2%; Pred. No. 9.9e-32;
 Matches 78; Conservative 14; Mismatches 22; Indels 2; Gaps 1;
 QY 3 GNDVQPPPTKICVCPDIPNPSPELEETLTHITKLAENNAFPFKIDNVKARVQV 62
 Db 252 GEDEL--PPMVCVCPKPIVDSPDLLEALNHSIAKLNAEHDGTFYFKIDTVKATVQV 309
 QY 63 AGKQYDFDVAERTCKSENELTESCKTKLGSLDCNAEVVYVPPWEKKLYPTV 118
 Db 310 GGLKYSIVFIARETTCKSNEELTKSCEINHGQILHCDNVYVPPWEKKYPTV 365
 RESULT 6
 KNG_MOUSE STANDARD; PRT; 661 AA.
 AC O08677; O08676; OS1XKS;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Kininogen precursor [Contains: Bradykinin].
 GN KNG.
 OS Mus musculus (Mouse).
 OC Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 [1]_TaxID=10090;
 RP SEQUENCE FROM N.A. (ISOPFORM HMW AND LMW).
 RC STRAIN=C57BL/6 X CBA; TISSUE=Liver;
 RX MEDLINE=97344355; PubMed=9199253;
 RA Takano M., Kondo J., Yayama K., Otani M., Sano K., Okamoto H.;
 RT "Molecular cloning of cDNAs for mouse low-molecular-weight and high-
 RT molecular-weight prekininogens."
 RL Biochim. Biophys. Acta 1352:222-230 (1997).
 RN [2]
 RP SEQUENCE FROM N.A. (ISOPFORM LMW).
 RC STRAIN=C57BL/6J; TISSUE=Placenta;
 RX MEDLINE=23354683; PubMed=12466851;
 RA Okazaki Y., Furuno M., Kasukawa T., Adachi J., Bono H., Kondo S.,
 RA Nakaide T., Osato N., Saito R., Suzuki H., Yamataka I., Kiyosawa H.,
 RA Yagi K., Tomaru Y., Hasegawa Y., Nogami A., Schonbach C., Gojibori T.,
 RA Baldairelli R., Hill D.P., Bult C., Rume D.A., Quackenbush J.,
 RA Schriber L.M., Kaspian A., Matsuda H., Batalov S., Beisel K.W.,
 RA Blake J.A., Bradt D., Brusic V., Chothia C., Corbani L.E., Cousins S.,
 RA Dalla E., Dragani T.A., Fletcher C.F., Forrest A., Frazer K.S.,
 RA Gaasterland T., Gariboldi M., Gissi C., Godzik A., Gough J.,
 RA Grimmond S., Gustincich S., Hirokawa N., Jackson I.J., Jarvis E.D.,
 RA Kanai A., Kawai H., Kawasawa Y., Kedzierski R.M., King B.L.,
 RA Konagaya A., Kurochkin I.V., Lee Y., Lenhard B., Lyons P.A.,
 RA Maglott D.R., Maltais L., Marchionni L., McKensie L., Miki H.,

RA Nageshima T., Numata K., Okido T., Pavan W.J., Pertea G., Pesole G.,
 RA Petrovsky N., Pillai R., Pontius J.U., Qi D., Ramachandran S.,
 RA Ravasi T., Reed J.C., Reed D.J., Reid J., Ring B.Z., Ringwald M.,
 RA Sandelin A., Schneider C., Sample C.A., Setou M., Shimada K.,
 RA Sultana R., Takenaka Y., Taylor M.S., Teasdale R.D., Tomita M.,
 RA Vitarolo R., Wagner L., Wahlestedt C., Wang Y., Watanabe Y., Wells C.,
 RA Wilmink L.G., Wymahaw-Boris A., Yanagisawa M., Yang I., Yang L.,
 RA Yuan Z., Zavalan M., Zhu Y., Zimmer A., Carninci P., Hayatsu N.,
 RA Hirozane-Kishikawa T., Konno H., Nakamura M., Sakazume N., Sato K.,
 RA Shiraki T., Waki K., Kawai J., Aizawa K., Arakawa T., Fukuda S.,
 RA Miyazaki A., Sakai K., Sasaki K., Shibata K., Shingawa A.,
 RA Yasunishi A., Yoshino M., Waterston R., Lander E.S., Rogers J.,
 RA Birney E., Hayashizaki Y.;
 RA "Analysis of the mouse transcriptome based on functional annotation of
 RT 60,770 full-length cDNAs."
 RL Nature 420:563-573 (2002).
 RW [3]
 RP SEQUENCE FROM N.A. (ISOPFORM LMW).
 RC TISSUE=Liver;
 RX MEDLINE=22388257; PubMed=12477932;
 RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.P., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.P., Casavant T.L., Scheetz T.E.,
 RA Raba S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullighy S.J.,
 RA Bonk S.A., McEwen P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay J.J., Ruliy S.W.,
 RA Villalon D.K., Murty D.N., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Rensy J., Heiton E., Kettman M., Madan A., Rodriguez S., Sanchez A.,
 RA Whitting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzywinski M.I., Skalek U., Smallos D.E.,
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length
 RT human and mouse cDNA sequences."
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
 CC -1- FUNCTION: (1) Kininogens are inhibitors of thiol proteases; (2)
 CC HMW-kininogen plays an important role in blood coagulation by
 CC helping to position optimally prekallikrein and factor XI next to
 CC factor XII; (3) HMW-kininogen inhibits the thrombin- and plasmin-
 CC induced aggregation of thrombocytes; (4) the active peptide
 CC bradykinin that is released from HMW-kininogen shows a variety of
 CC physiological effects: (4A) influence in smooth muscle
 CC contraction, (4B) induction of hypotension, (4C) natriuresis and
 CC diuresis, (4D) decrease in blood glucose level, (4E) it is a
 CC mediator of inflammation and causes (4E1) increase in vascular
 CC permeability, (4E2) stimulation of nociceptors (4E3) release of
 CC other mediators of inflammation (e.g. prostaglandins), (4F) it has
 CC a cardioprotective effect (directly via bradykinin action),
 CC indirectly via endothelium-derived relaxing factor action; (5)
 CC LMW-kininogen inhibits the aggregation of thrombocytes; (6) LMW-
 CC kininogen is in contrast to HMW-kininogen not involved in blood
 CC clotting (by similarity).
 CC -1- SUBCELLULAR LOCATION: Secreted.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Name=HMW;
 CC IsoId=O08677-1; Sequence=Displayed;
 CC Name=LMW;
 CC IsoId=O08677-2; Sequence=VSP_001263; VSP_001264;
 CC -1- TISSUE SPECIFICITY: Plasma.
 CC -1- PTM: Bradykinin is released from kininogen by plasma kallikrein.
 CC -1- SIMILARITY: Contains 3 cystatin-like domains.
 CC -----
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DR EMBL, M11984; AAA41487.1; ..
DR EMBL, M14369; AAA41484.1; ..
DR EMBL, M14369; AAA41485.1; ALT_SEQ.
DR EMBL, M16455; AAA41482.1; ..
DR PIR, A25486; A25486.
DR PIR, A28055; A28055.
DR InterPro; IPR000010; Cystatin.
DR InterPro; IPR002395; Kininogen.
DR Pfam; PF00031; cystatin; 3.
DR PRINTS; PR00334; KININOGEN.
DR SMART; SM00043; CY; 3.
DR PROSITE; PS00287; CYSTATIN; 2.
KW Glycoprotein; Plasma; Repeat; Thiol protease inhibitor; Vasodilator;
KW Bradykinin; Blood coagulation; Inflammatory response; Signal,
KW Alternative splicing; Multigene family.
FT SIGNAL 1 18
FT CHAIN 19 389 KININOGEN.
FT CHAIN 19 380 KININOGEN HEAVY CHAIN.
FT PEPTIDE 381 389 BRADYKININ.
FT CHAIN 390 639 KININOGEN LIGHT CHAIN.
FT DOMAIN 19 136 CYSTATIN-LIKE 1.
FT DOMAIN 137 258 CYSTATIN-LIKE 2.
FT DOMAIN 259 380 CYSTATIN-LIKE 3.
FT DOMAIN 439 514 HIS-RICH.
FT DISULFID 28 609 INTERCHAIN (BY SIMILARITY).
FT DISULFID 83 94 BY SIMILARITY.
FT DISULFID 107 126 BY SIMILARITY.
FT DISULFID 142 145 BY SIMILARITY.
FT DISULFID 206 218 BY SIMILARITY.
FT DISULFID 229 248 BY SIMILARITY.
FT DISULFID 264 267 BY SIMILARITY.
FT DISULFID 328 340 BY SIMILARITY.
FT DISULFID 351 370 BY SIMILARITY.
FT CARBOHYD 82 82 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 127 127 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 169 169 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 205 205 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 294 294 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 529 529 N-LINKED (GLCNAC. .) (POTENTIAL).
FT VARSPLIC 400 433 VSPVIAVQVERDPGNEQPIHGHMLHAKO -> RLNS
FT VARSPLIC 400 433 CEYKGRLLKAGAPAPERQAEASTVTP (in isoform
FT VARSPLIC 400 433 LMW)
FT VARSPLIC 434 639 /FTid-vsp 001265.
FT CONFLICT 61 61 Missing (in isoform LMW).
FT SEQUENCE 639 AA; 70933 MW; D3172DF94FF56AP5 CRC64;
Query Match 59.7%; Score 410; DB 1; Length 639;
Best Local Similarity 66.4%; Pred. NO. 2e-31;
Matches 77; Conservative 13; Mismatches 26; Indels 0; Gaps 0;
OY 3 GKDPVPTKICVCPDRPTNSPELETLTHITKLNAENATFVKIDNVKARQV 62
DB 253 GDLFLPDLPCPCPRNPVPSPELKEALGHSIAQLAENHTFFKIDTVKATSV 312
OY 63 AGKVPYDFVARETTCKSENEETSCETKLGSLDCHAEVTVVPEKKIYPTV 118
DB 313 AGTKVIEPIARETKCKSENAELTADCETKRLGSLNCNANVYRPNKVVPTV 368
STANDARD; PRT: 430 AA.
RESULT 8
KNT2_RAT
ID KNT2_RAT
AC P08932;
DT 01-NOV-1988 (Rel. 09, Created)
DT 01-NOV-1988 (Rel. 09, Last sequence update)
DT 15-MAR-2004 (Rel. 43, Last annotation update)
DE T-kininogen II precursor (Major acute phase protein) (Alpha-1-MAP)
DE (Tiosastatin) [contains: T-kinin].
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

```

```

OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=86008264; PubMed=2413018;
RA Furuto-Kato S., Matsumoto A., Kitamura N., Nakanishi S.;
RT "Primary structures of the mRNAs encoding the rat precursors for
RT bradykinin and T-kinin. Structural relationship of kininogens with
RT major acute phase protein and alpha 1-cysteine proteinase
RT inhibitor.";
RL J. Biol. Chem. 260:12054-12059(1985).
CC -!- FUNCTION: Kininogens are plasma glycoproteins with a number of
CC functions: (1) as precursor of the active peptide bradykinin they
CC effect smooth muscle contraction, induction of hypotension and
CC increase of vascular permeability. (2) They play a role in blood
CC coagulation by helping to position optimally prekallikrein and
CC factor XI next to factor XII. (3) They are inhibitor of thiol
CC proteases.
CC -!- SUBCELLULAR LOCATION: Extracellular.
CC -!- TISSUE SPECIFICITY: Plasma.
CC -!- INDUCTION: In response to an inflammatory stimulant. T-kininogen
CC II synthesis is induced and the plasma concentration of
CC T-kininogen I is raised.
CC -!- PTM: AS T-KININ IS PRECEDED BY A MET INSTEAD OF AN ARG OR LYS, IT
CC IS NOT RELEASED FROM ITS PRECURSOR BY EITHER TISSUE OR PLASMA
CC KALLIKREIN.
CC -!- MISCELLANEOUS: Rats express four types of kininogens: the
CC classical HMW and LMW kininogens produced by alternative splicing
CC of the same gene, and two additional LMW-like kininogens: T-I and
CC T-II.
CC -!- SIMILARITY: Contains 3 cystatin-like domains.
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CC entities requires a license agreement (See http://www.ebi.ac.uk/announcements
CC or send an email to license@ebi.ac.uk).
EMBL, M11895; AAA41491.1; ..
PIR, B28055; B28055.
DR GlycoSuiteDB; G08932; ..
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; cystatin; 3.
DR SMART; SM00043; CY; 3.
DR PROSITE; PS00287; CYSTATIN; 2.
KW Glycoprotein; Plasma; Repeat; Vasodilator; Multigene family;
KW Thiol protease inhibitor; Bradykinin; Acute phase; Signal.
FT SIGNAL 1 18
FT CHAIN 19 430 KININOGEN, T-II.
FT CHAIN 19 375 HEAVY CHAIN.
FT PEPTIDE 376 386 LIGHT CHAIN.
FT CHAIN 387 430 CYSTATIN-LIKE 1.
FT DOMAIN 19 135 CYSTATIN-LIKE 2.
FT DOMAIN 136 257 CYSTATIN-LIKE 3.
FT DOMAIN 258 375 INTERCHAIN (BY SIMILARITY).
FT DISULFID 28 404 BY SIMILARITY.
FT DISULFID 83 94 BY SIMILARITY.
FT DISULFID 107 125 BY SIMILARITY.
FT DISULFID 141 144 BY SIMILARITY.
FT DISULFID 205 217 BY SIMILARITY.
FT DISULFID 228 247 BY SIMILARITY.
FT DISULFID 263 266 BY SIMILARITY.
FT DISULFID 327 339 BY SIMILARITY.
FT DISULFID 350 369 BY SIMILARITY.
FT CARBOHYD 82 82 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 126 126 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 168 168 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 204 204 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 326 326 N-LINKED (GLCNAC. .) (POTENTIAL).
SQ SEQUENCE 430 AA; 47524 MW; 43EDF02D1BF55076 CRC64;
Query Match 56.5%; Score 388; DB 1; Length 430;

```

Best Local Similarity 62.1%, Pred. No. 1.6e-29, Matches 72; Conservative 15; Mismatches 29; Indels 0; Gaps 0;	
QY	3 GKDVPQPTKICVGGCPDIPNTPNSELSTLTHITKLNANNATPFIKIDNVKARQVV 62
Db	252 GDDLPKLLPKKFCGPKIPVDSPLKALGHSAQLNAQHLPFKIDTVKATGVV 311
QY	63 AGKYFIDVARETTCSENEELTSCETKULGSLDGNASVYVPEKKIYPTV 118
Db	312 AGTKYVIEPIARETNCQNTLTDCEYKLGSLDGNANVYRPENKVVPTV 367
RESULT 9	
IDENT1 RAT	STANDARD; PRT; 430 AA.
AC	P01046; P04081;
DT	01-NOV-1986 (Rel. 03, Created)
DT	01-NOV-1988 (Rel. 09, Last sequence update)
DT	15-MAR-2004 (Rel. 43, Last annotation update)
DE	7-kininogen I precursor (Major acute phase protein) (Alpha-1-MAP)
DE	(Thioesterin) [Contains: T-kinin].
GN	MAP1.
OS	Rattus norvegicus (Rat).
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC	Mammalia; Eutharia; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX	NCBI_TaxID=10116;
RN	[1]
RP	SEQUENCE FROM N.A.
RA	MEDLINE=86008264; PubMed=2413018;
RA	Purito-Kato S., Matsumoto A., Kitamura N., Nakanishi S.;
RT	"Primary structures of the mRNAs encoding the rat precursors for
RT	bradykinin and T-kinin. Structural relationship of kininogens with
RT	major acute phase protein and alpha 1-cysteine proteinase
RT	inhibitor."
RT	J. Biol. Chem. 260:12054-12059 (1985).
RN	[2]
RP	SEQUENCE OF "5-430 FROM N.A., AND PARTIAL SEQUENCE.
RA	MEDLINE=86008266; PubMed=2413019;
RA	Anderson K.P., Heath E.C.;
RT	"The relationship between rat major acute phase protein and the
RT	kininogens."
RT	J. Biol. Chem. 260:12065-12071 (1985).
RN	[3]
RP	SEQUENCE OF 7-430 FROM N.A.
RA	MEDLINE=85127561; PubMed=2578992;
RA	Cole I., Inglis A.S., Roxburgh C.M., Howlett G.J., Schreiber G.;
RT	"Major acute phase alpha 1-protein of the rat is homologous to bovine
RT	kininogen and contains the sequence for bradykinin: its synthesis is
RT	regulated at the mRNA level."
RT	FEBS Lett. 182:57-61 (1985).
RN	[4]
RP	SEQUENCE OF 1-65 FROM N.A.
RA	MEDLINE=87250580; PubMed=2439509;
RA	Pung W.-P., Schreiber G.;
RT	"Structure and expression of the genes for major acute phase alpha 1-
RT	protein (thioesterin) and kininogen in the rat."
RT	J. Biol. Chem. 262:9298-9308 (1987).
CC	-1- FUNCTION: Kininogens are plasma glycoproteins with a number of
CC	functions: (1) as precursor of the active peptide bradykinin they
CC	effect smooth muscle contraction, induction of hypotension and
CC	increase of vascular permeability. (2) They play a role in blood
CC	coagulation by helping to position optimally prekallikrein and
CC	factor XI next to factor XII. (3) They are inhibitor of thiol
CC	proteases.
CC	-1- SUBCELLULAR LOCATION: Extracellular.
CC	-1- TISSUE SPECIFICITY: Plasma.
CC	-1- INDUCTION: In response to an inflammatory stimulant. T-kininogen
CC	II synthesis is induced and the plasma concentration of
CC	T-kininogen I is raised.
CC	-1- PTM: AS T-KININ IS PRECEDED BY A MET INSTEAD OF AN ARG OR LYS, IT
CC	IS NOT RELEASED FROM ITS PRECURSOR BY EITHER TISSUE OR PLASMA
CC	KALLIKREIN.
CC	-1- MISCELLANEOUS: Rate express four types of kininogens: the

CC	classical HMW and LMW kininogens produced by alternative splicing
CC	of the same gene, and two additional LMW-like kininogens: T-I and
CC	T-II.
CC	-1- SIMILARITY: Contains 3 cystatin-like domains.
CC	-1- CAUTION: In addition to the conflicts described in the feature
CC	table, Ref.2 sequence differs from that shown in positions 257,
CC	262, 268, 269, 295, 314, 315, 331, 332 and 389. In all those
CC	positions the alternate amino acid is the one present in T-II
CC	kininogens.
CC	-----
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CC	between the Swiss Institute of Bioinformatics and the EMBL outstation -
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CC	use by non-profit institutions as long as its content is in no way
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CC	entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC	or send an email to license@isb-sib.ch).
CC	-----
CC	EMBL; M11893; AAA41489.1; ..
CC	EMBL; M1661; AAA41570.1; ..
CC	EMBL; M16454; AAA41568.1; ..
CC	EMBL; X02299; CAA26162.1; ALT_SEQ.
CC	PIR; A01286; KGRIT1.
CC	PIR; A21897; A23897.
CC	PIR; A27115; A27115.
CC	GlycosuiteDB; P01046; ..
CC	Interpro; IPR000010; Cystatin.
CC	PIfam; PF00031; cystatin; 3.
CC	SMART; PS00043; CY; 3.
CC	PROSITE; PS00287; CYSTATIN; 2.
CC	Glycoprotein; Plasma; Repeat; Vasodilator; Multigene family;
CC	Thiol protease inhibitor; Bradykinin; Acute phase; Signal.
CC	SIGNAL 1 18
CC	CHAIN 19 430 KININOGEN, T-I.
CC	HEAVY CHAIN.
CC	PEPTIDE 376 386
CC	CHAIN 387 430
CC	DOMAIN 19 135
CC	DOMAIN 136 257
CC	DOMAIN 258 375
CC	DISULFID 28 404
CC	DISULFID 83 94
CC	DISULFID 107 125
CC	DISULFID 141 144
CC	DISULFID 205 217
CC	DISULFID 228 247
CC	DISULFID 263 266
CC	DISULFID 327 339
CC	DISULFID 350 369
CC	CARBOHYD 82 82
CC	CARBOHYD 126 126
CC	CARBOHYD 168 168
CC	CARBOHYD 204 204
CC	CARBOHYD 326 326
CC	CONFLICT 26 28
CC	CONFLICT 55 55
CC	CONFLICT 61 61
CC	CONFLICT 83 83
CC	CONFLICT 166 166
CC	CONFLICT 179 181
CC	CONFLICT 193 193
CC	CONFLICT 212 212
CC	CONFLICT 214 214
CC	CONFLICT 229 229
CC	CONFLICT 233 233
CC	CONFLICT 257 257
CC	CONFLICT 262 262
CC	CONFLICT 264 264
CC	CONFLICT 268 269
CC	CONFLICT 295 295
CC	CONFLICT 314 315
CC	CONFLICT 331 332
CC	CONFLICT 389 389

PT CONFLICT 414 414 R -> G (IN REF. 2 AND 3).
PT CONFLICT 415 415 A -> L (IN REF. 2).
PT CONFLICT 420 421 DH -> ER (IN REF. 3).
PT CONFLICT 430 430 P -> S (IN REF. 1).
SQ SEQUENCE 430 AA, 47715 MW, PAEBB78FAF4723C3 CRC64;

Query Match 55.3%, Score 380, DB 1, Length 430;
Best Local Similarity 62.1%, Pred No. 98-29, 30, Indels 0, Gaps 0;
Matches 72, Conservative 14, Mismatches 30, Indels 0, Gaps 0;

OY 3 GDFVQPTKICVCCPRDPTNSPELEETLTHTITKLAENNAATPYFKINVKARVQV 62
DB 252 GDFVQPTKICVCCPRDPTNSPELEETLTHTITKLAENNAATPYFKINVKARVQV 62

OY 63 AGKYPIDFVARETTCSKESBELTESCETKKGSLDCAEAVTVVWEKKIYFTV 118
DB 312 AGVIVIEFIARETNCQSKQTELTADCTKHGSLDCAEAVTVVWEKKIYFTV 367

RESULT 10
ID_CYTF_MOUSE STANDARD, PRT, 144 AA.
AC 089058;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Cystatin P precursor (Leukocystatin) (Cystatin 7) (Cystatin-like
DE metastasis-associated protein) (CMAP).
GN CST7.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98299157, PubMed=9632704;
RA Halfon S., Ford J., Foster J., Dowling L., Lucian L., Sterling M.,
RA Xu Y., Weiss M., Ikeda M., Ligggett D., Helms A., Caux C., Lebecque S.,
RA Hannum C., Menon S., McClanahan T., Gorman D., Zurawski G.,
RA "Leukocystatin, a new class II cystatin expressed selectively by
RA hematopoietic cells".
RL J. Biol. Chem. 273:16400-16408(1998).
CC -!- FUNCTION: Inhibits pepsin and cathepsin L but with affinities
CC lower than other cystatins. May play a role in immune regulation
CC through inhibition of a unique target in the hematopoietic system.
CC -!- SUBCELLULAR LOCATION: Secreted (Probable).
CC -!- SIMILARITY: Belongs to the cystatin family.

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CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
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CC entities requires a license agreement (See <http://www.ebi.ac.uk/announcements/>
CC or send an email to license@ebi.ac.uk).

DR EMBL; AF031826; AAC40140.1;
DR EMBL; AF031825; AAC40139.1;
DR HSPG; P01034; IG96.
DR MGD; MGI:1298217; Cst7.
DR InterPro; IPR000010; Cystatin.
DR Pfam; PF00031; Cystatin; 1.
DR SMART; SM00043; Cst1.
DR PROSITE; PS00287; CYSTATIN; FALSE NEG.
KW Thiol protease inhibitor; Glycoprotein; Signal.
FT SIGNAL 1 18
FT CHAIN 19 144
FT ACT_SITE 36 36
FT SITE 80 84
FT DISULFID 98 109
FT DISULFID 123 143
FT SEQUENCE 144 AA; 16380 MW; B5837334C1B4A89C CRC64;

Query Match 25.0%, Score 171.5, DB 1, Length 144;
Best Local Similarity 35.5%, Pred No. 1-7e-09;
Matches 39, Conservative 22, Mismatches 42, Indels 7, Gaps 3;

OY 4 KDFVQPTKICVCCPRDPTNSPELEETLTHTITKLAENNAATPYFKINVKARVQV 63
DB 27 KDLI---SSVKGPFPTTETNNPGLKAARHSVEKNNCTNDIPLFKSHVSKALVQV 83

OY 64 GKYPIDFVARETTCSKESBELTESCETKKGSLDCAEAVTVVWEKKIYFTV 110
DB 84 GLKYPLEVIGRTTCKTMMHQL-DNCDPQTNPALAKETLYCYSEVVVPI 132

RESULT 11
ID_CYTF_HUMAN STANDARD, PRT, 145 AA.
AC 076056; Q9UED4;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 10-OCT-2003 (Rel. 42, Last annotation update)
DE Cystatin P precursor (Leukocystatin) (Cystatin 7) (Cystatin-like
DE metastasis-associated protein) (CMAP).
GN CST7.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98406131, PubMed=9733783;
RA Ni J., Fernandez M.A., Daniellson L., Chillakuru R.A., Zhang J.,
RA Grubb A., Su J., Gentz R., Abrahamson M.,
RA "Cystatin P is a glycosylated human low molecular weight cysteine
RA proteinase inhibitor".
RL J. Biol. Chem. 273:24797-24804(1998).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=98299157, PubMed=9632704;
RA Halfon S., Ford J., Foster J., Dowling L., Lucian L., Sterling M.,
RA Xu Y., Weiss M., Ikeda M., Ligggett D., Helms A., Caux C., Lebecque S.,
RA Hannum C., Menon S., McClanahan T., Gorman D., Zurawski G.,
RA "Leukocystatin, a new class II cystatin expressed selectively by
RA hematopoietic cells".
RL J. Biol. Chem. 273:16400-16408(1998).
RN [3]
RP SEQUENCE FROM N.A.
RA Morita M., Arakawa H., Yoshiuchi N.;
RA "Human homologue of murine CMAP".
RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE FROM N.A.
RX MEDLINE=20399571, PubMed=10945474;
RA Morita M., Hara Y., Tanai Y., Arakawa H., Nishimura S.;
RA "Genomic construct and mapping of the gene for CMAP
RA (Leukocystatin/Cystatin P, CST7) and identification of a proximal
RA novel gene, BSCV (C20orf3)".
RL Genomics 67:87-91(2000).
RN [5]
RP SEQUENCE FROM N.A.
RX MEDLINE=21638749, PubMed=11780052;
RA Deloukas P., Matthews L.H., Ashurst J., Burton J., Gilbert J.G.R.,
RA Jones M., Stavrides G., Almeida J.P., Babbage A.K., Baguley C.L.,
RA Bailey J., Barlow K.P., Bates K.N., Beard L.M., Beare D.M.,
RA Beasley O.P., Bird C.P., Blakey S.B., Bridgman A.M., Brown A.J.,
RA Buck D., Burrill W.D., Butler A.P., Carder C., Carter N.P.,
RA Chapman J.C., Clamp M., Clark G., Clark L.N., Clark S.Y., Clee C.M.,
RA Clegg S., Cobley V.B., Collier R.E., Connor R.B., Corby N.R.,
RA Coulson A., Coville G.J., Deadman R., Dhami P.D., Dunn M.R.,
RA Ellington A.G., Frankland J.A., Fraser A., French L., Garner P.,
RA Grafham D.V., Griffiths C., Griffiths M.N.D., Gwilliam R., Hall R.E.,
RA Hammond S., Harley J.B., Heath P.D., Ho S., Holden J.L., Howden P.J.,
RA Huckle E., Hunt A.R., Hunt S.E., Jackson K., Johnson C.M., Johnson D.,
RA Kay M.P., Kimberley A.M., King A., Knightes A., Laird G.K., Lawlor S.,

Lehvaeslahti M.H., Leversha M.A., Lloyd C., Lloyd D.M., Lovell J.D., Marsh V.B., Martin S.L., McConachie I.J., McLeay K., McMurray A.A., Milne S.A., Mistry D., Moore M.J.P., Mullikin J.C., Nickerson T., Oliver K., Parker A., Patel R., Pearce T.A.V., Peck A.I., Phillips B.J.C.T., Prathalingam S.R., Plumb R.W., Ramsay H., Rice C.M., Ross M.T., Scott C.B., Sehra H.K., Showkhen R., Sims S., Skuce C.D., Smith M.T., Soderlund C., Stewart C.A., Sulston J.B., Swann R.M., Sycamore N., Taylor R., Tee L., Thomas D.W., Thorpe A., Tracey A., Tromans A.C., Vaudin M., Wall M., Wallis J.M., Whitehead S.L., Whittaker P., Willey D.L., Williams L., Williams S.A., Wilming L., Wray P.W., Hubbard T., Durbin R.M., Bentley D.R., Beck S., Rogers J., Nature 414:865-871 (2001).

SEQUENCE FROM N.A.

RC TISSUE=lung;

RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G., Klausner R.D., Collins P.S., Wagner L., Shenmen C.M., Schuler G.D., Altschul S.F., Zdobych B., Buetow K.H., Schaefer C.P., Bhat N.K., Hopkins R.P., Jordan H., Moore T., Max S.I., Wang J., Hsieh F., Datchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L., Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E., Brownstein M.J., Udwin T.B., Toshiyuki S., Carninci P., Prange C., Reha S.S., Lequellano N.A., Peters G.J., Abramson R.D., Mullahy S.J., Bonak S.A., McSwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H., Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Rulyk S.W., Villalón D.K., Muny D.M., Sodergren E.J., Lu X., Gibbs R.A., Pahey J., Hailton E., Kettman N., Madan A.C., Rodriguez S., Sanchez A., Blakeley R.W., Touchman J.W., Green E.D., Dickson M.C., Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butterfield Y.S.N., Krzywinski M.I., Skaleka U., Smalhus D.E., Schnerch A., Schein J.E., Jones S.J.M., Marra M.A., "Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences."

RT PROC. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).

RL FUNCTION: Inhibits papain and cathepsin L but with affinities lower than other cystatins. May play a role in immune regulation through inhibition of a unique target in the hematopoietic system.

CC SUBCELLULAR LOCATION: Secreted (Probable).

CC TISSUE SPECIFICITY: Primarily expressed in peripheral blood cells and spleen.

CC SIMILARITY: Belongs to the cystatin family.

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DR EMBL: AF036342; AAC35747.1; .

DR EMBL: AF031824; AAC39768.1; .

DR EMBL: AB015225; BAB34941.1; ALT_INIT.

DR EMBL: AB029636; BAB11866.1; ALT_INIT.

DR EMBL: AL035661; CAB75498.1; .

DR EMBL: BC015507; AAH15507.1; ALT_INIT.

DR HSSP: P01034; 1G96.

DR Genbank: HGNC:2479; CST7.

DR MIM: 603253; .

DR GO: GO:0004869; P:cysteine protease inhibitor activity; TAS.

DR GO: GO:0006955; P:immune response; TAS.

DR InterPro: IPR000010; Cystatin.

DR SMART: SM00031; cystatin; 1.

DR Pfam: SM00043; CY; 1.

DR PROSITE: PS00287; CYSTATIN; 1.

DR Thiol protease inhibitor; Anyloids; Signal.

FT SIGNAL 1 19

FT CHAIN 20 145

FT ACT_SITE 37 37

REACTIVE SITS.

FT SITE 81 85 SECONDARY AREA OF CONTACT.

FT DISULFID 99 110 BY SIMILARITY.

FT DISULFID 124 144 BY SIMILARITY.

FT CARBOHYD 62 62 N-LINKED (GLCNAC...) (POTENTIAL).

FT CARBOHYD 115 115 N-LINKED (GLCNAC...) (POTENTIAL).

SO SEQUENCE 145 AA; 16454 MW; B2BC4P76857CB0F CRC64;

Query Match 33.8%; Score 163.5; DB 1; Length 145;

Best Local Similarity 31.6%; Pred. No. 9.9e-09;

Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

QY 11 TKICVGPDPPTNSPELETHITIKLAENNAFTYFKIDNVKKARVQVQVAKKVFID 70

DB 32 SRVKGPPKTIKTNDPGVLAARYSEKFNCTNDMFLKESRITRALVQIVGLKYMLE 91

QY 71 FVARETTCSKESNEELTESCE---TKKQSLDCNAEYVVPWBKKIYPTVTYVHWE 124

DB 92 VEIGRTTCKQKHRL-DCDFQTNHTLQTLSCYSEVWVWVW-----LQHFE 138

RESULT 12

CYT_CMACMU STANDARD; PRT; 146 AA.

AC O19092;

DT 15-JUL-1998 (Rel. 36, Created)

DT 15-JUL-1998 (Rel. 36, Last sequence update)

DT 28-FEB-2003 (Rel. 41, Last annotation update)

DE Cystatin C precursor.

GN CST3.

OS Macaca mulatta (Rhesus macaque).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea; OC Cercopithecoidea; Macaca.

OX NCBI_Taxid=95544;

RN Cystatin C precursor.

RP MEDLINE=97054523; PubMed=8988820;

RA Wei L.H., Walker L.C., Levy E.;

RT "Cystatin C, Icelandic-like mutation in an animal model of cerebrovascular beta-amyloidosis."

RL Stroke 27:2080-2085 (1996).

CC FUNCTION: As an inhibitor of cysteine proteinases, this protein is thought to serve an important physiological role as a local regulator of this enzyme activity.

CC SIMILARITY: Belongs to the cystatin family.

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DR EMBL: U51912; AAB64050.1; .

DR HSSP: P01034; 1G96.

DR InterPro: IPR000010; Cystatin.

DR Pfam: PF00031; Cystatin; 1.

DR SMART: SM00043; CY; 1.

DR PROSITE: PS00287; CYSTATIN; 1.

DR Thiol protease inhibitor; Anyloids; Signal.

FT SIGNAL 1 26

FT CHAIN 27 146

FT ACT_SITE 37 37

FT SITE 81 85

FT DISULFID 99 109

FT DISULFID 123 143

SO SEQUENCE 146 AA; 15857 MW; F0B3BB774A39DF26 CRC64;

Query Match 20.2%; Score 138.5; DB 1; Length 146;

Best Local Similarity 27.9%; Pred. No. 2.4e-06;

Matches 34; Conservative 25; Mismatches 52; Indels 11; Gaps 4;

QY 8 OPTKICVCGPRDIPNTPSPLESTLTHITKLNENATFYFKIDNVKQVQVAGKY 67
 Db 31 KPFR--LVGPMDSVZEEGRVAFSEVTKNSNDMYSRALQVVRARKQVAGVNY 88
 QY 68 FIDFVARTTCKESNELTESC---ETKLGSLDNCNAEVVVPVWKKIYPTVTWNHWE 124
 Db 89 FLVDELGTCTK--TOPNLDNCPPEOPHLKAKAFCSFOIYTVFMO----GWTLSKST 142
 QY 125 CE 126
 Db 143 CQ 144

RESULT 13

CYTM_HUMAN
 ID CYTM_HUMAN STANDARD; PRT; 149 AA.
 AC Q15628;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 15-MAR-2004 (Rel. 43, Last annotation update)
 DE Cystatin M precursor (Cystatin E).
 GN C8T6.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN (1)
 RP SEQUENCE FROM N.A.
 RX MEDLINE=97150844; PubMed=8995380;
 RA Sotiropoulos G., Anisowicz A., Sager R.;
 RT "Identification, cloning, and characterization of cystatin M, a novel
 RT cysteine proteinase inhibitor, down-regulated in breast cancer.";
 RL J. Biol. Chem. 272:903-910(1997).
 RN (2)
 RP SEQUENCE FROM N.A.
 RX MEDLINE=97256812; PubMed=9099741;
 RA Ni J., Abrahamson M., Zhang M., Fernandez M.A., Grubb A., Su J.,
 RA Yu G.L., Li Y., Parmelee D., Xing L., Coleman T.A., Gentz S.,
 RA Thakura R., Nguyen N., Hesselberg M., Gentz R.;
 RT "Cystatin E is a novel human cysteine proteinase inhibitor with
 RT structural resemblance to family 2 cystatins.";
 RL J. Biol. Chem. 272:10853-10858(1997).
 RN (3)
 RP SEQUENCE FROM N.A.
 RX TISSUE=Prostate;
 RX MEDLINE=2238257; PubMed=12477932;
 RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins P.S., Wagner L., Shenman C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Rabe S.S., Loughran N.A., Peters G.J., Abramson R.D., Mullen S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S.S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Faney J., Helton E., Kettner M., Madan A., Rodriguez S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzywinski M.I., Skalska U., Smallos D.E.,
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length
 RT human and mouse cDNA sequences."
 RN Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
 RN (4)
 RP CHARACTERIZATION, AND TISSUE SPECIFICITY.
 RX MEDLINE=21246880; PubMed=1114845;
 RA Zeewen P.L., Van Vlijmen-Willems I.M., Jansen B.J., Sotiropoulos G.,
 RA Curfs J.H., Mels J.F., Jansen J.J., Van Kuissen P., Schalkwijk J.;
 RT "Cystatin M/E expression is restricted to differentiated epidermal
 RT keratinocytes and sweat glands: a new skin-specific proteinase

RT inhibitor that is a target for cross-linking by transglutaminase.";
 RL J. Invest. Dermatol. 116:693-701(2001).
 CC - FUNCTION: Shows moderate inhibition of cathepsin B but is not
 CC active against cathepsin C.
 CC - SUBCELLULAR LOCATION: Secreted.
 CC - TISSUE SPECIFICITY: Restricted to the stratum granulosum of normal
 CC skin, the stratum granulosum/spinosum of psoriatic skin, and the
 CC secretory coils of eccrine sweat glands. Low expression levels are
 CC found in the nasal cavity.
 CC - PFM: Substrate for transglutaminases. Acts as an acyl acceptor but
 CC not as an acyl donor.
 CC - SIMILARITY: Belongs to the cystatin family.
 CC
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 CC
 CC EMBL; U62800; RAB06566.1;
 DR EMBL; U81233; RAB06566.1;
 DR EMBL; BC011334; RAB06566.1;
 DR HSRP; P01038; ICEW.
 DR Gene; HGNC:2478; CST6.
 DR MIM; 601891;
 DR GO; GO:0004869; F:cysteine protease inhibitor activity; TAS.
 DR GO; GO:0007345; P:embryogenesis and morphogenesis; TAS.
 DR InterPro; IPR000010; Cystatin.
 DR Pfam; PF00031; cystatin; 1.
 DR SMART; SM00043; CV; 1.
 DR PROSITE; PS00287; CYSTATIN; 1.
 KW SIGNAL protease inhibitor; Signal, Glycoprotein.
 FT SIGNAL 1 28
 FT CHAIN 29 149
 FT ACT SITE 26 36
 FT SITE 26 36
 FT DISULFID 98 113
 FT DISULFID 126 146
 FT CARBOHYD 137 137
 SQ SEQUENCE 149 AA; 16511 MW; 2076A78BFC9FAC8C CRC64;
 Query Match 20.2%; Score 138.5; DB 1; Length 149;
 Best Local Similarity 31.5%; Pred. No. 2.4e-06;
 Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;
 QY 10 PTKICVCGPRDIPNTPSPLESTLTHITKLNENATFYFKIDNVKQVQVAGKYFI 69
 Db 30 PFRVWGLRLDLPDDQVOKAAQAVSYNGMSITFRDTHIIKQSLVAGIKYFL 89
 QY 70 DFVARETTCSE-----SNEELTESCTYKLGQ--SLDCNAEVVVPVWE 111
 Db 90 TREMGSTDCRTRVTDGVDLT-TCPLAAGAQQKLCDFEVLVVPVQ 136
 RESULT 14
 CYTC_BOVIN
 ID CYTC_BOVIN STANDARD; PRT; 148 AA.
 AC P01035;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 15-JUL-1999 (Rel. 35, Last sequence update)
 DT 28-FEB-2003 (Rel. 43, Last annotation update)
 DE Cystatin C precursor (Colostrum thiol proteinase inhibitor).
 GN C8T3.
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovidae; Bovinae; Bos.
 OX NCBI_TaxID=9913;
 RN (1)
 RP SEQUENCE FROM N.A.; SEQUENCE OF 66-83, AND CHARACTERIZATION.
 RN TISSUE=Cerebrospinal fluid, and Choroid plexus;

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RX MEDLINE=98094199; PubMed=9434110;
RA Olsson S.-L., Ek B., Wilm M., Broberg S., Raak L., Björk I.,
RT Molecular cloning and N-terminal analysis of bovine cystatin C
RT Identification of a full-length N-terminal region."
RL Biochim. Biophys. Acta 1343:203-210(1997).
RN [2]
RP SEQUENCE OF 37-148.
RX MEDLINE=85311205; PubMed=3991407;
RA Hirado M., Tsunawake S., Sakiyama F., Minobe M., Fujii S.;
RT "Complete amino acid sequence of bovine colostrum low-Mr cysteine
RT proteinase inhibitor."
RL FEBS Lett. 186:41-45(1985).
RX MEDLINE=85311205; PubMed=3991407;
RA Hirado M., Tsunawake S., Sakiyama F., Minobe M., Fujii S.;
RT "Complete amino acid sequence of bovine colostrum low-Mr cysteine
RT proteinase inhibitor."
RL FEBS Lett. 186:41-45(1985).
CC -1- FUNCTION: This is a thiol proteinase inhibitor.
CC -1- MASS SPECTROMETRY: MW=13420; METHOD=MALDI.
CC -1- SIMILARITY: Belongs to the cystatin family.
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CC -----
CC EMBL; Y10811; CAA71771.1; -.
CC HSP; P01034; I96.
CC InterPro; IPR000010; Cystatin.
CC Pfam; PF00031; cystatin; 1.
CC SMART; SM00043; CY; 1.
CC PROSITE; PS00287; CYSTATIN; 1.
KM Thiol protease inhibitor; signal; Pyroglutamate carboxylic acid.
FT SIGNAL 1 30
FT CHAIN 31 148
FT MOD_RES 31 31 PYROGLUTAMATE CARBOXYLIC ACID (PROBABLE).
FT ACT_SITE 40 40 REACTIVE SITE.
FT SITE 84 88 SECONDARY AREA OF CONTACT.
FT DISULFID 102 112 BY SIMILARITY.
FT DISULFID 126 146 BY SIMILARITY.
SQ SEQUENCE 148 AA; 16265 MW; BE740FE37CEB9FOE CRC64;
Query Match 20.0%; Score 137.5; DB 1; Length 148;
Best Local Similarity 28.8%; Pred. No. 3e-06;
Matches 32; Conservative 25; Mismatches 35; Indels 19; Gaps 4;
Oy 24 NSPELSETHITKLNENATFYKIDNVKQVGVAGKYFIDFVARETTCSKSN 83
Db 48 NEEGVQELSFVAVSEFKRSDAVQSRVVRVARKQVGMVYFLDVELGRTYTK--S 105
Oy 84 EELTBSG-----STKGLQSLDCNAEVVYVPEKKIYPTVTVNHWCE 126
Db 106 QANLDSCPFHNPQHLKREK-----CSFQVYVVPWN-----TINLVKFSQ 147
RESULT 15
PETS_RAT
ID_PETS_RAT STANDARD; PRT; 378 AA.
AC Q9QX79;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Fetuin-B precursor (IRL685).
GN FETUB.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Eutelestomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Sprague-Dawley; TISSUE=Liver;
RX MEDLINE=20407138; PubMed=10947975;
RA Olivier E., Soury E., Ruminy P., Rouson A., Parmentier P., Davenau M.,
RA Saller J.-P.;
RT "Fetuin-B, a second member of the fetuin family in mammals."

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RL Biochem. J. 350:589-597(2000).
CC -1- SUBCELLULAR LOCATION: Secreted (Potential).
CC -1- TISSUE SPECIFICITY: Liver.
CC -1- SIMILARITY: Belongs to the fetuin family.
CC -1- SIMILARITY: Contains 2 cystatin-like domains.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; AJ242926; CAB62543.1; -.
CC InterPro; IPR000010; Cystatin.
CC Pfam; PF00031; cystatin; 2.
CC SMART; SM00043; CY; 2.
CC PROSITE; PS01254; FETUIN_1; 1.
CC PROSITE; PS01255; FETUIN_2; 1.
KM Glycoprotein; signal; Repeat.
FT SIGNAL 1 18
FT CHAIN 19 378
FT DOMAIN 27 152
FT DOMAIN 153 273
FT DISULFID 96 107
FT DISULFID 120 140
FT DISULFID 154 157
FT DISULFID 217 224
FT DISULFID 237 260
FT CARBOHYD 40 40 N-LINKED (GLCNAC...) (POTENTIAL).
FT CARBOHYD 139 139 N-LINKED (GLCNAC...) (POTENTIAL).
SQ SEQUENCE 378 AA; 41532 MW; 066C0A5C3B03C878 CRC64;
Query Match 19.7%; Score 135; DB 1; Length 378;
Best Local Similarity 25.8%; Pred. No. 1.5e-05;
Matches 33; Conservative 32; Mismatches 49; Indels 14; Gaps 5;
Oy 7 VQPPTK-----ICVGCPRDPTNSPSELTHTITKLNENATFYKIDNVKQV 61
Db 142 LRPVSKRKIHSMCDPCPHVDLSAPVLEAATSLAKFNSENPSKQYALV-KVTKATQW 200
Oy 62 VAGKYFIDFVARETTCSKSNELTESCTKGLQSLDCNAEVVYVPEKKIYPTV 119
Db 201 VVGPSPYFVEYLKESPTQSDSCSLQASDSEPVGL---COGSLKSPGVFPQPKKTVT 257
Oy 120 VNHWECEP 127
Db 258 VS---CEP 262
Search completed: September 24, 2004, 14:09:13
Job time : 9.636 secs

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R; Kitamura, N.; Kitagawa, H.; Fukushima, D.; Takagaki, Y.; Miyata, T.; Nakanishi, S.
 J. Biol. Chem. 260, 8610-8617, 1985
 A:Title: Structural organization of the human kininogen gene and a model for its evolution
 A:Reference number: A2545; MUID:85234583; PMID:2989294
 A:Contents: annotation; gene organization
 R; Pierce, J.V.
 Fed. Proc. 27, 52-57, 1968
 A:Title: Structural features of plasma kinins and kininogens.
 A:Reference number: A31455; MUID:50255622; PMID:4952632
 A:Contents: annotation; bradykinin
 C:Comment: The LMW kininogen precursor is produced from the same gene as the HMW form (a)
 C:Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
 C:Comment: bradykinin, released from kininogen by kallikrein, is a potent vasodilator, and
 C:Comment: xiprolin residue is present in the kininogen prior to the release of bradykinin.
 C:Genetics:
 A:Gene: GDB:KNG
 A:Cross-references: GDB:125256; OMIM:228960
 A:Map position: 3q27-3q27
 A:Introns: 65/3; 102/3; 131/1; 188/3; 224/3; 253/1; 310/3; 346/3; 375/3; 401/3
 C:Superfamily: kininogen; cystatin homology
 C:Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; glyco
 P:1-18/Domain: signal sequence #status predicted <SIG>
 P:19-427/Product: LMW prokininogen (kininogen I) #status predicted <KAT>
 P:19-389/Product: LMW kininogen II #status predicted <KAT>
 P:19-379/Product: LMW kininogen heavy chain #status predicted <KAT>
 P:19-331/Domain: cystatin homology <CY>
 P:142-253/Domain: cystatin homology <CY2>
 P:264-375/Domain: cystatin homology <CY3>
 P:380-385/Product: lysyl-bradykinin (kallidin I) #status experimental <KBDY>
 P:381-389/Product: bradykinin (kallidin I) #status experimental <BDY>
 P:390-427/Product: LMW kininogen light chain #status experimental <LCH>
 P:128-407/Modified site: pyroglutamate carboxylic acid (Gln) (in mature form) #status predicted
 P:148-169/205,294/Binding site: carboxylate (Asn) (covalent) #status predicted
 P:379-380/Cleavage site: Met-Lys (kallikrein) #status experimental
 P:383/Modified site: 4-hydroxyproline (pro) (partially) #status experimental
 P:389-390/Cleavage site: Arg-Ser (kallikrein) #status experimental
 P:401/Binding site: carboxylate (Thr) (covalent) #status absent

Query Match 90.0%; Score 618; DB 1; Length 427;
 Best Local Similarity 100.0%; Pred. No. 3.6e-50;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY	3	GKDFVQPTKICVGCPRDPTNSPELETLTHITIKLNAENNAATFYFKIDNVKARVQV 62
DB	253	GKDFVQPTKICVGCPRDPTNSPELETLTHITIKLNAENNAATFYFKIDNVKARVQV 312
QY	63	AGKYPTDFVARETTCKESNELTESCETKLGQSLDCNAEYVVPWEKKIYPTV 118
DB	313	AGKYPTDFVARETTCKESNELTESCETKLGQSLDCNAEYVVPWEKKIYPTV 368

RESULT 2
 KGNH1
 N:Alcarnate names: alpha-2-thiol proteinase inhibitor - human
 N:Contains: bradykinin (kallidin I); HMW kininogen I; HMW kininogen II; low molecular we
 C:Species: Homo sapiens (man)
 C:Date: 28-May-1986 #sequence revision 28-May-1986 #text change 08-Dec-2000
 A:Accession: A01279; A25276; S32422; A91153; A24871; A27699; A31905; A34030; S02
 R:Ohkubo, I.; Kurachi, K.; Takagawa, T.; Shiohara, H.; Sasaki, M.
 Biochemistry 23, 5691-5697, 1984
 A:Title: Isolation of a human cDNA for alpha-2-thiol proteinase inhibitor and its identi
 A:Reference number: A90490; MUID:85122621; PMID:6441591
 A:Accession: A01279
 A:Molecule type: mRNA
 A:Residues: 1-389 <CHK>
 A:Cross-references: GDB:K05566; NID:9177889
 R: Takagaki, Y.; Kitamura, N.; Nakanishi, S.
 J. Biol. Chem. 260, 8601-8609, 1985
 A:Title: Cloning and sequence analysis of cDNAs for human high molecular weight and low
 A:Reference number: A92544; MUID:85234582; PMID:2989293
 A:Accession: A25276

A:Molecule type: mRNA
 A:Residues: 1-592; '1', 594-644 <TAK>
 A:Cross-references: GB:M11437; NID:9386852
 R: Auerwald, E.A.; Roessler, D.; Mentale, R.; Aebfeld-Machleidt, I.
 FEBS Lett. 321, 93-97, 1993
 A:Title: Cloning expression and characterization of human kininogen domain 3.
 A:Reference number: S32422; MUID:93223854; PMID:8467516
 A:Accession: S32422
 A:Molecule type: mRNA
 A:Residues: 'ANSW', 353-377 <AUE>
 A:Note: differences are due to known cloning artifacts
 R: Lottspeich, F.; Kellermann, J.; Henschen, A.; Foerster, B.; Muller-Esterl, W.
 Eur. J. Biochem. 153, 307-314, 1985
 A:Title: The amino acid sequence of the light chain of human high-molecular-mass kininogen
 A:Reference number: A91153; MUID:86030270; PMID:4054110
 A:Accession: A91153
 A:Molecule type: protein
 A:Residues: 379-644 <LOT>
 A:Note: the bradykinin sequence preceding the light chain sequence was not determined in
 R: Kellermann, J.; Lottspeich, F.; Henschen, A.; Muller-Esterl, W.
 Eur. J. Biochem. 153, 471-478, 1986
 A:Title: Completion of the primary structure of human high-molecular-mass kininogen. The
 A:Reference number: A24871; MUID:86108361; PMID:3484703
 A:Accession: A24871
 A:Molecule type: protein
 A:Residues: '2', 20-380 <KEL1>
 R: Kellermann, J.; Lottspeich, F.; Henschen, A.; Muller-Esterl, W.
 in Kinins IV, Greenbaum, L.M., and Margolis, H.S., ed., pp.85-89, Plenum Press, New York
 A:Title: Amino acid sequence of the light chain of human high molecular mass kininogen.
 A:Reference number: A27699; MUID:88209021; PMID:3365237
 A:Accession: A27699
 A:Molecule type: protein
 A:Residues: 380-389 <MIN>
 R: Maeda, H.; Matsumura, Y.; Kato, H.
 J. Biol. Chem. 263, 16051-16054, 1988
 A:Title: Purification and identification of [hydroxyprolyl(3)]bradykinin in ascitic fluid
 A:Reference number: A31905; MUID:89034061; PMID:3182782
 A:Accession: A31905
 A:Molecule type: protein
 A:Residues: 381-389 <KAR>
 R: Saaguri, M.; Ikeda, M.; Ideishi, M.; Arakawa, K.
 Biochem. Biophys. Res. Commun. 150, 511-516, 1988
 A:Title: Identification of [hydroxyproline(3)]-lysyl-bradykinin released from human plasma
 A:Reference number: A34030; MUID:88106632; PMID:3337729
 A:Accession: A34030
 A:Molecule type: protein
 A:Residues: 380-389 <SAS>
 R: Lenarcic, B.; Gabrijelcic, D.; Rozman, B.; Drobnic-Kosorok, M.; Turk, V.
 Biol. Chem. Hoppe-Seyler 369, 257-261, 1988
 A:Title: Human cathelain B and cysteine proteinase inhibitors (CPIs) in inflammatory and
 A:Reference number: S02482; MUID:89076517; PMID:3264507
 A:Accession: S02482
 A:Molecule type: protein
 A:Residues: 310-314, 381-389 <LEN1>
 R: Kato, H.; Matsumura, Y.; Maeda, H.
 FEBS Lett. 232, 252-254, 1988
 A:Title: Isolation and identification of hydroxyproline analogues of bradykinin in human
 A:Reference number: A61495; MUID:88211869; PMID:3366244
 A:Accession: A61495
 A:Molecule type: protein
 A:Residues: 380-389 <KAT1>
 A:Experimental source: urine
 A:Note: this peptide had Pro-383 modified to 4-hydroxyproline
 A:Accession: B61495
 A:Molecule type: protein
 A:Residues: 381-389 <KAT2>

A:Experimental source: urine
A:Note: this peptide had Pro-383 modified to 4-hydroxyproline
A:Accession: C61495
A:Molecule type: Protein
R:Lenardic, B.; Krasovec, M.; Ritonja, A.; Olafsson, I.; Turk, V.
FEBS Lett. 280, 211-215, 1991
A:Title: Inactivation of human cystatin C and kininogen by human cathepsin D.
A:Reference number: S14303; MUID:9119133; PMID:2013314
A:Accession: S14447
A:Molecule type: Protein
A:Residues: 364-359, 'N', 361-375 <LEN2>
R:Little, S.S.; Johnson, D.A.
Biochem. J. 307, 341-346, 1995
A:Title: Human mast cell tryptase isoforms: separation and examination of substrate-specificity
A:Reference number: S55239; MUID:95251593; PMID:7733667
A:Accession: S55239
A:Molecule type: Protein
A:Residues: 450-452, 'X', 454, 'X', 456 <LIT>
R:Straczek, J.; Maschi, F.; Le Nguyen, D.; Becchi, M.; Heulin, M.H.; Nebet, P.; Bellevil
FEBS Lett. 373, 207-211, 1995
A:Title: Purification from human plasma of a tetrapeptide that potentiates insulin-like
A:Reference number: S68059; MUID:96033974; PMID:7589467
A:Accession: S68059
A:Molecule type: Protein
A:Residues: 431-434 <GRA>
R:Kitamura, N.; Kitagawa, H.; Fukushima, D.; Takagaki, Y.; Miyata, T.; Nakanishi, S.
J. Biol. Chem. 260, 8610-8617, 1985
A:Title: Structural organization of the human kininogen gene and a model for its evolution
A:Reference number: A9548; MUID:85234583; PMID:2989394
A:Contents: annotation; gene organization
R:Pierce, J.V.
Fed. Proc. 27, 52-57, 1968
A:Title: Structural features of plasma kinins and kininogens.
A:Reference number: A91455; MUID:90255622; PMID:4952632
A:Contents: annotation; bradykinin
C:Comment: The HMW kininogen precursor and the LMW form are produced from the same gene
C:Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
C:Comment: The glycine/histidine/lysine-rich region of HMW kininogen light chain is important
C:Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, a
C:Comment: xypoline residue is present in the kininogen prior to the release of bradykinin.
C:Genetics:
A:Gene: GDB:KNG
A:Cross-references: GDB:125256; OMIM:228960
A:Map position: 3q27-3q27
A:Introns: 65/3; 102/3; 131/1; 188/3; 224/3; 253/1; 310/3; 346/3; 375/3
C:Superfamily: kininogen; cystatin homology
C:Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; dupl
F:19-644/Product: HMW kininogen I (prokininogen) #status experimental <MAT1>
F:19-379,390-644/Product: HMW kininogen II #status experimental <MAT2>
F:19-379/Domains: HMW kininogen heavy chain #status experimental <HCH>
F:19-131/Domains: cystatin homology <CY1>
F:142-253/Domains: cystatin homology <CY2>
F:264-375/Domains: cystatin homology <CY3>
F:380-389/Product: lysyl-bradykinin (kallidin II) #status experimental <KBDY>
F:381-389/Product: bradykinin (kallidin I) #status experimental <BDY>
F:390-644/Domains: HMW kininogen light chain #status experimental <LCH>
F:431-434/Region: glycine/histidine/lysine-rich 30-residue repeats
F:431-434/Product: low molecular weight growth promoting factor #status experimental <GF>
F:19/Modified site: pyroglutamate carboxylic acid (Gln) (in mature form) #status experimental
F:28-614,83-94,107-126,142-145,206-218,229-248,264-267,328-340,351-370/Disulfide bonds:
F:48/Binding site: carboxylate (Asn) (covalent) #status experimental
F:169,205,294/Binding site: carboxylate (Asn) (covalent) #status experimental
F:379-380/Cleavage site: Met-Lys (kallikrein) (partial) #status experimental
F:383/Modified site: 4-hydroxyproline (Pro) #status experimental
F:389-390/Cleavage site: Arg-Ser (kallikrein) #status experimental
F:401,533,542,546,557,571,593,628/Binding site: carboxylate (Thr) (covalent) #status ex
F:577/Binding site: carboxylate (Ser) (covalent) #status experimental
Query Match 90.0%; Score 618; DB 1; Length 644;
Best Local Similarity 100.0%; Pred. No. 5.7e-50;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVGCPRDIPNTSPLESTLTHTITKLNANNATFYFKIDNVKARQVW 62
DB 253 GKDFVQPTKICVGCPRDIPNTSPLESTLTHTITKLNANNATFYFKIDNVKARQVW 312
QY 63 AGKKYPTDFVARETTCKSNBELTESCETKLGSLDCNAEVVVPWEKKIYPTV 118
DB 313 AGKKYPTDFVARETTCKSNBELTESCETKLGSLDCNAEVVVPWEKKIYPTV 368
RESULT 3
KGBOL1
N:Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen
N:Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen
C:Species: Bos primigenius taurus (cattle)
C:Date: 14-Nov-1983 #sequence_revision 14-Nov-1983 #text_change 22-Jun-1999
C:Accession: A01283
R:Nawa, H.; Kitamura, N.; Hirose, T.; Arai, M.; Inayama, S.; Nakanishi, S.
Proc. Natl. Acad. Sci. U.S.A. 80, 90-94, 1983
A:Title: Primary structures of bovine liver low molecular weight kininogen precursors
A:Reference number: A93984; MUID:83117859; PMID:6572010
A:Accession: A01283
A:Molecule type: mRNA
A:Residues: 1-436 <NWA>
A:Cross-references: GB:J00010; GB:V00426; NID:G163256; PID:AAA30604.1; PID:G163257
C:Comment: The LMW kininogen precursor is produced from the same gene as the HMW form
C:Comment: Kininogen is a cysteine proteinase inhibitor, takes part in initiation of
C:Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator
C:Comment: xypoline residue is present in the kininogen prior to the release of bradykinin.
C:Superfamily: kininogen; cystatin homology
C:Keywords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; g
F:1-18/Domains: signal sequence #status predicted <SIG>
F:18-436/Product: LMW kininogen I #status predicted <MAT>
F:18-378/Product: LMW kininogen I heavy chain #status predicted <HCH>
F:18-130/Domains: cystatin homology <CY1>
F:141-252/Domains: cystatin homology <CY2>
F:263-374/Domains: cystatin homology <CY3>
F:379-388/Product: lysyl-bradykinin (kallidin II) #status predicted <KBDY>
F:380-388/Product: bradykinin (kallidin I) #status predicted <BDY>
F:389-436/Product: LMW kininogen I light chain #status experimental <LCH>
F:19/Modified site: pyroglutamate carboxylic acid (Gln) (in mature form) #status predic
F:27-406,82-93,106-125,141-144,205-217,228-248,263-266,327-339,350-369/Disulfide bond
F:47,87,168,169,197,204/Binding site: carboxylate (Asn) (covalent) #status predicted
F:378-379/Cleavage site: Met-Lys (kallikrein) #status predicted
F:382/Modified site: 4-hydroxyproline (Pro) #status predicted
F:388-389/Cleavage site: Arg-Ser (kallikrein) #status predicted
Query Match 64.0%; Score 440; DB 1; Length 436;
Best Local Similarity 70.4%; Pred. No. 1.7e-33;
Matches 81; Conservative 14; Mismatches 20; Indels 0; Gaps 0;
QY 4 KDFVQPTKICVGCPRDIPNTSPLESTLTHTITKLNANNATFYFKIDNVKARQVW 63
DB 253 KDFVQPTKICVGCPRDIPNTSPLESTLTHTITKLNANNATFYFKIDNVKARQVW 312
QY 64 GKKYPTDFVARETTCKSNBELTESCETKLGSLDCNAEVVVPWEKKIYPTV 118
DB 313 GKKYPTDFVARETTCKSNBELTESCETKLGSLDCNAEVVVPWEKKIYPTV 367
RESULT 4
KGBOL1
N:Alternate names: alpha-2-thiol proteinase inhibitor; preprokininogen
N:Contains: bradykinin (kallidin); kininogen I; kininogen II; prokininogen
C:Species: Bos primigenius taurus (cattle)
C:Date: 14-Nov-1983 #sequence_revision 14-Nov-1983 #text_change 22-Jun-1999
C:Accession: A01281; A91923; A91938; A29559
R:Kitamura, N.; Takagaki, Y.; Furuto, S.; Tanaka, T.; Nawa, H.; Nakanishi, S.
Nature 305, 545-549, 1983
A:Title: A single gene for bovine high molecular weight and low molecular weight kin
A:Reference number: A93317; MUID:84014106; PMID:6571899

J. Biochem. 67, 313-323, 1970
 A>Title: Studies on the structure of bovine kininogen: cleavage of disulfide bonds and
 A/Reference number: A91923; MUID:70180420; PMID:4986212
 A/Accession: A91923
 A/Molecule type: Protein
 A/Residues: 376-391 <Mat>
 R/Han, Y.N.; Kato, H.; Iwanaga, S.; Suzuki, T.
 J. Biochem. 79, 1201-1222, 1976
 A>Title: Primary structure of bovine plasma high-molecular-weight kininogen. The amino
 A/Reference number: A91941; MUID:76260155; PMID:956151
 A/Accession: A91941
 A/Molecule type: Protein
 A/Residues: 387-455 <HAN>
 A/Note: 398-Pro, 401-Val, and 455-Lys were also found
 R/Han, Y.N.; Komiya, M.; Iwanaga, S.; Suzuki, T.
 J. Biochem. 77, 55-68, 1975
 A>Title: Studies on the primary structure of bovine high-molecular-weight kininogen. Amino
 A/Reference number: A91938; MUID:75170265; PMID:1169237
 A/Accession: A91938
 A/Molecule type: Protein
 A/Residues: 436-496 <H2>
 R/Sueyoshi, T.; Miyata, T.; Hashimoto, N.; Kato, H.; Hayashida, H.; Miyata, T.; Iwanaga, S.
 J. Biol. Chem. 262, 2768-2779, 1987
 A>Title: Bovine high molecular weight kininogen. The amino acid sequence, positions of
 A/Reference number: A92627; MUID:87137530; PMID:3546295
 A/Accession: B29859
 A/Molecule type: Protein
 A/Residues: 2, 20-104, 106-256, 257-376 <SUS>
 R/Lottepeich, P.; Kallermann, J.; Henschen, A.; Foerster, B.; Muller-Esterl, W.
 Eur. J. Biochem. 152, 307-314, 1985
 A>Title: The amino acid sequence of the light chain of human high-molecular-mass kininogen
 A/Reference number: A91153; MUID:86030270; PMID:4054110
 A/Contents: annotation; bovine cleavage sites; bovine carbohydrate binding sites
 R/Sueyoshi, T.; Miyata, T.; Kato, H.; Iwanaga, S.
 Seikagaku 56, 808, 1984
 A>Title: Disulfide bonds in bovine HMW kininogens.
 A/Reference number: A94300
 A/Contents: annotation; disulfide bonds
 A/Note: article in Japanese
 C/Comment: The HMW kininogen precursor is produced from the same gene as the LMW form as
 C/Comment: kininogen is a cysteine proteinase inhibitor, takes part in initiation of the
 C/Comment: The glycine/histidine/lysine-rich region of HMW kininogen light chain is imprecisely
 C/Comment: Bradykinin, released from kininogen by kallikrein, is a potent vasodilator, and
 C/Comment: xiprolone residue is present in the kininogen prior to the release of bradykinin.
 C/Superfamily: kininogen; cystatin homology
 C/KeyWords: alternative splicing; blood coagulation; cysteine proteinase inhibitor; duplication
 F/19-619/Product: HMW kininogen II #status predicted <SIG>
 F/19-376/Product: HMW kininogen II #status predicted <MAT>
 F/19-130/Product: HMW kininogen II heavy chain #status experimental <HC>
 F/141-252/Domain: cystatin homology <CY1>
 F/1261-372/Domain: cystatin homology <CY2>
 F/377-386/Product: lysyl-bradykinin (kallidin II) #status experimental <MBDY>
 F/378-386/Product: bradykinin (kallidin I) #status experimental <BDY>
 F/387-619/Product: HMW kininogen II light chain #status experimental <LCH>
 F/418-488/Region: glycine/histidine/lysine-rich
 F/19/Modified site: pyrrolidone carboxylic acid (Gln) (in mature form) #status experimental
 F/27-589, 82-93, 106-125, 141-144, 205-217, 228-247, 261-264, 325-337, 348-367/Disulfide bonds:
 F/47/Binding site: carbohydrate (Asn) (covalent) #status absent
 F/167, 168, 169, 204, 280/Binding site: carbohydrate (Asn) (covalent) #status experimental
 F/136/Binding site: carbohydrate (Thr) (covalent) (partial) #status experimental
 F/197/Binding site: carbohydrate (Asn) (covalent) (partial) #status experimental
 F/376-377/Cleavage site: Met-Lys (kallikrein) #status experimental
 F/380/Modified site: 4-hydroxyproline (pro) #status predicted
 F/386-387/Cleavage site: Arg-Ser (kallikrein) #status experimental
 F/396, 400, 404, 510/Binding site: carbohydrate (Ser) (covalent) #status experimental
 F/397, 398, 518, 522, 534, 546, 551, 569/Binding site: carbohydrate (Thr) (covalent) #status experimental
 F/496-497/Cleavage site: Arg-Thr (kallikrein) #status experimental

Query Match 60.1%; Score 413; DB 1; Length 619;
 Best Local Similarity 67.2%; Pred. No. 8, 28-31;
 Matches 78; Conservative 14; Mismatches 22; Indels 2; Gaps 1;

QY 3 GKDFVQPTKICVGCPRDIPNTPSPLESLTHTITKLAENNATFYFKIDNVKARVQV 62
 DB 252 GEDFL--PPMVCGCPKPIVDSPDLAEALNHSIAKLNAEHDGTFYFKIDNVKARVQV 309
 QY 63 AGKKYFIDPVARETTCSENEELTSSCTKKLGQSLDCNARVYVVPWEKKIYPTV 118
 DB 310 GGLKYSIVPIARETTCSENEELTSSCTKKLGQSLDCNARVYVVPWEKKIYPTV 365

RESULT 7

A28055
 K:kininogen, LMW I precursor - rat
 C:Species: Rattus norvegicus (Norway rat)
 C/Date: 20-Jun-1989 #sequence_revision 20-Jun-1989 #text_change 15-Nov-1996
 C/Accession: A28055
 R:Purito-Kato, S.; Matsumoto, A.; Kitamura, N.; Nakanishi, S.
 J. Biol. Chem. 260, 12054-12059, 1985
 A>Title: Primary structures of the mRNAs encoding the rat precursors for bradykinin and
 inhibitor.
 A/Reference number: A92496; MUID:86008264; PMID:2413018
 A/Accession: A28055
 A/Molecule type: mRNA
 A/Residues: 1-433 <PUR>
 C/Superfamily: kininogen; cystatin homology
 C/KeyWords: alternative splicing
 F/1-18/Domain: signal sequence #status predicted <SIG>
 F/19-433/Product: K-kininogen, LMW I #status predicted <MAT>
 F/19-131/Domain: cystatin homology <CY1>
 F/142-253/Domain: cystatin homology <CY2>
 F/264-375/Domain: cystatin homology <CY3>

Query Match 59.7%; Score 410; DB 2; Length 433;
 Best Local Similarity 66.4%; Pred. No. 1, 1e-30;
 Matches 77; Conservative 13; Mismatches 26; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVGCPRDIPNTPSPLESLTHTITKLAENNATFYFKIDNVKARVQV 62
 DB 253 GDDLPELLSPDCPCPNIPVDSPELKEALGHSIAQLNENHTFYFKIDNVKATSOV 312

QY 63 AGKKYFIDPVARETTCSENEELTSSCTKKLGQSLDCNARVYVVPWEKKIYPTV 118
 DB 313 AGTKYVIEPIARETTCSENEELTSSCTKKLGQSLDCNARVYVVPWEKKIYPTV 368

RESULT 8

A25486
 K:kininogen, HMW I precursor - rat
 N:Contains: bradykinin
 C:Species: Rattus norvegicus (Norway rat)
 C/Date: 08-Mar-1989 #sequence_revision 08-Mar-1989 #text_change 15-Nov-1996
 R/Kitagawa, H.; Kitamura, N.; Hayashida, H.; Miyata, T.; Nakanishi, S.
 J. Biol. Chem. 262, 2190-2198, 1987
 A>Title: Differing expression patterns and evolution of the rat kininogen gene family.
 A/Reference number: A92625; MUID:87137443; PMID:3029068
 A/Accession: A25486
 A/Molecule type: mRNA
 A/Residues: 1-639 <KIT>
 A/Note: the authors translated the codon CAA for residue 347 as Aen
 C/Superfamily: kininogen; cystatin homology
 C/KeyWords: alternative splicing
 F/1-18/Domain: signal sequence #status predicted <SIG>
 F/19-639/Product: kininogen, HMW I #status predicted <MAT>
 F/19-131/Domain: cystatin homology <CY1>
 F/142-253/Domain: cystatin homology <CY2>
 F/264-375/Domain: cystatin homology <CY3>

Query Match 59.7%; Score 410; DB 2; Length 639;
 Best Local Similarity 66.4%; Pred. No. 1, 6e-30;
 Matches 77; Conservative 13; Mismatches 26; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVGCPRDIPNTPSPLESLTHTITKLAENNATFYFKIDNVKARVQV 62
 DB 252 GEDFL--PPMVCGCPKPIVDSPDLAEALNHSIAKLNAEHDGTFYFKIDNVKARVQV 309
 QY 63 AGKKYFIDPVARETTCSENEELTSSCTKKLGQSLDCNARVYVVPWEKKIYPTV 118
 DB 310 GGLKYSIVPIARETTCSENEELTSSCTKKLGQSLDCNARVYVVPWEKKIYPTV 365

R; Furuto-Kato, S.; Matsumoto, A.; Kitamura, N.; Nakanishi, S.
 J. Biol. Chem. 260, 12054-12059, 1985
 A; Title: Primary structures of the mRNAs encoding the rat precursors for bradykinin and inhibitor.
 A; Reference number: A92496; MUID:86008264; PMID:2413018
 A; Accession: A01286
 A; Molecule type: mRNA
 A; Residues: 1-430 <RUR>
 A; Cross-references: GB:M11883, NID:G205084; PID:G205085
 R; Kitagawa, H.; Kitamura, N.; Hayashida, H.; Miyata, T.; Nakanishi, S.
 J. Biol. Chem. 262, 2190-2198, 1987
 A; Title: Differing expression patterns and evolution of the rat kininogen gene family.
 A; Reference number: A92625; MUID:87137443; PMID:3029068
 A; Accession: D25486
 A; Molecule type: DNA
 A; Residues: 375-430 <KIT>
 R; Enjiyoi, K.; Kato, H.; Hayashi, I.; Oh-ishi, S.; Iwanaga, S.
 J. Biol. Chem. 263, 973-979, 1988
 A; Title: Purification and characterization of rat T-kininogens isolated from plasma of a rabbit.
 A; Reference number: A92729; MUID:88087226; PMID:3121623
 A; Accession: A28526
 A; Molecule type: protein
 A; Residues: E'20-48; 376-430 <ENJ>
 R; Xanda, S.; Sugiyama, K.; Takahashi, M.; Shumiyu, S.; Tomino, S.; Nagase, S.
 Jpn. J. Cancer Res. 81, 63-68, 1990
 A; Title: Identification of a protein increasing in serum of Nagase analbuminemic rats by immunoblotting.
 A; Reference number: PLO193; MUID:90216390; PMID:2108948
 A; Accession: PLO193
 A; Molecule type: mRNA
 A; Residues: 330-420, 'P', '422-429, 'P', <KAN>
 R; Anderson, K.P.; Croyle, M.L.; Lingrel, J.B.
 Gene 81, 119-128, 1989
 A; Title: Primary structure of a gene encoding rat T-kininogen.
 A; Reference number: JQ0027; MUID:90034172; PMID:2806908
 A; Accession: JQ0027
 A; Molecule type: DNA
 A; Residues: 1-60, 'E', '62-113, 'R', '115-165, 'P', '167-178, 'TKI', '182-211, 'P', '213-256, 'S', '258-388 <KAN>
 R; Experimental source: Strain Sprague-Dawley
 R; Kagayama, R.; Kitamura, N.; Ohkubo, H.; Nakanishi, S.
 J. Biol. Chem. 262, 2345-2351, 1987
 A; Title: Differing utilization of homologous transcription initiation sites of rat K and X kininogen.
 A; Reference number: A25488; MUID:87137465; PMID:3818598
 A; Accession: B25488
 A; Status: preliminary
 A; Molecule type: DNA
 A; Residues: 1-48 <KAG>
 A; Cross-references: GB:M14356, NID:G205090; PID:AAA41492.1; PID:G205091
 R; Enjiyoi, K.; Kato, H.; Hayashi, I.; Oh-ishi, S.; Iwanaga, S.
 J. Biol. Chem. 263, 965-972, 1988
 A; Title: Purification and characterization of two kinds of low molecular weight kininogen.
 A; Reference number: A28525; MUID:88087225; PMID:3335530
 A; Accession: A28525
 A; Molecule type: protein
 A; Residues: 376-430 <EN2>
 R; Sierra, F.; Walter, R.; Vautravers, P.; Guigoz, Y.
 Arch. Biochem. Biophys. 322, 333-338, 1995
 A; Title: Identification of several isoforms of T-kininogen expressed in the liver of aged rats.
 A; Reference number: S68034; MUID:96032652; PMID:7574705
 A; Accession: S68036
 A; Molecule type: mRNA
 A; Residues: 340-430 <SIS>
 A; Experimental source: clone pSG17
 A; Comment: At least three types of LMW kininogen precursors are present in rat plasma, coding bradykinin.
 C; Comment: T-kininogens contain T-kinin (I-S-bradykinin), a novel kinin isolated after digestion of an Arg or Lys, it is probably not released from its precursor by either tissue or R.
 C; Comment: The T-kininogens are produced in response to an inflammatory stimulant.
 C; Genes: 65/3, 102/3, 130/1, 187/3, 223/2, 252/1, 309/3, 345/3, 374/3, 398/3
 C; Superfamily: kininogen; cystatin homology
 C; Keywords: acute phase; bradykinin; cysteine proteinase inhibitor; duplication; glycoprotein
 F; 11-18/Domain: signal sequence #status predicted <SIG>
 F; 19-430/Product: T-kininogen I #status experimental <MAT>

F; 19-130/Domain: cystatin homology <CY1>
 F; 141-252/Domain: cystatin homology <CY2>
 F; 263-374/Domain: cystatin homology <CY3>
 F; 378-386/Product: bradykinin #status predicted <BDY>
 F; 19/Modified site: pyroglutamate carboxylic acid (Gln) (in mature form) #status expirime
 F; 82, 126, 168, 204, 326/Binding site: carboxylate (Asn) (covalent) #status predicted
 F; 83-94, 107-125, 141-144, 205-217, 228-247, 263-266, 327-339, 350-369/Disulfide bonds: #statu
 Query Match 55.3%, Score 380; DB 1; Length 430;
 Best Local Similarity 62.1%, Pred. No. 6.7e-28;
 Matches 72; Conservative 14; Mismatches 30; Indels 0; Gaps 0;
 QY 3 GDFVQPTKLCVQCDPTPTSPLESLTITITKLNENNATPFYKIDNVKARVQV 62
 DB 252 GDLFLPLNKGCPREIPVDSPKLGALGHSAGLNAQHNFYFKITVAKASQV 311
 QY 63 AKKYPIDFVARETTCKSNBELTESCTKKLGSLDCNAEVVVPWEKKIYPTV 118
 DB 312 AGVIIVIEFTARETNCOSKSTELTADCTHGLGSLNCNANVYMRPWKNVPTV 367
 RESULT 13
 UDBO
 Cystatin - bovine
 N; Alternate names: thiol proteinase inhibitor
 C; Species: Bos primigenius taurus (cattle)
 C; Date: 28-Feb-1986 #sequence_revision 28-Feb-1986 #text_change 06-Dec-1996
 C; Accession: A01271
 R; Hirado, M.; Teunissen, S.; Sakiyama, P.; Ninobe, M.; Fujii, S.
 FEBS Lett. 186, 41-45, 1985
 A; Title: Complete amino acid sequence of bovine colostrum low-M-r cysteine proteinase I
 A; Reference number: A01271; MUID:85231205; PMID:3891407
 A; Accession: A01271
 A; Molecule type: protein
 A; Residues: 1-112 <HIR>
 C; Superfamily: cystatin; cystatin homology
 C; Keywords: colostrum; cysteine proteinase inhibitor
 F; 2-112/Domain: cystatin homology <CY>
 F; 48-52/Region: inhibitory #status predicted
 F; 66-76, 90-110/Dissulfide bonds: #status predicted
 Query Match 20.0%, Score 137.5; DB 1; Length 112;
 Best Local Similarity 28.8%, Pred. No. 7.5e-06;
 Matches 32; Conservative 25; Mismatches 35; Indels 19; Gaps 4;
 QY 24 NSPELEETLTITKLNENNATPFYKIDNVKARVQVAGKYPIDFVARETTCKSN 83
 DB 12 NEEGVQALSPAVGEFNKRNDAVQGRVVRVVRARQVVGNYPLDVELGRTTCTK--S 69
 QY 84 ESLTESC-----ETKLGSLDCNAEVVVPWEKKIYPTVTVHWECE 126
 DB 70 QANLSCFPNQLKREKL-----CSFQVYVVPWVN---TINLVKFSQ 111
 RESULT 14
 UDBO
 Cystatin C precursor [validated] - human
 N; Alternate names: gamma-CSP; gamma-trace; neuroendocrine basic polypeptide; post-gamm
 C; Species: Homo sapiens (man)
 C; Date: 06-Jul-1982 #sequence_revision 31-Mar-1991 #text_change 08-Dec-2000
 C; Accession: S10216; S00004; J10095; A33400; S02751; A01270; A25434; S12288; A32732; A
 R; Abrahamson, M.; Olafsson, I.; Palsdottir, A.; Ulvabæk, M.; Lundwall, A.; Jensen, J
 Biochem. J. 268, 287-294, 1990
 A; Title: Structure and expression of the human cystatin C gene.
 A; Reference number: S10216; MUID:90303202; PMID:2363674
 A; Accession: S10216
 A; Molecule type: DNA
 A; Residues: 1-146 <ABL>
 A; Cross-references: EMBL:X5255; NID:G30257; PID:CAA36497.1; PID:G296643
 R; Abrahamson, M.; Grubb, A.; Olafsson, I.; Lundwall, A.
 FEBS Lett. 216, 229-233, 1987
 A; Title: Molecular cloning and sequence analysis of cDNA coding for the precursor of cl
 A; Reference number: S00004; MUID:87219149; PMID:3495457

A:Residues: 8-49 <ESN>
 R:Enard, A.; Enard, F.; Guillou, F.; Gauthier, F.
 FEBS Lett. 300, 131-135, 1992
 A:Title: Production of the cysteine proteinase inhibitor cystatin C by rat Sertoli cells
 A:Reference number: S21109, MUID:92225121, PMID:1563513
 A:Accession: S21109

A:Molecule type: protein
 A:Residues: 8, 'XX', 11-20 <ES2>
 C:Superfamily: cystatin; cystatin homology
 C:Keywords: cysteine proteinase inhibitor
 F:15-127/Domain: cystatin homology <CYS>
 F:80-90,104-124/Disulfide bonds: #status predicted

Query Match 18.9%; Score 130; DB 2; Length 127;
 Best Local Similarity 28.0%; Pred. No. 4.3e-05;
 Matches 30; Conservative 28; Mismatches 43; Indels 6; Gaps 4;

Oy	8	QPTKICVGPDRDIPNTPSELEETLTHITKLNAENNATFYKIDNVKARVQVYVAGKY	67
Db	11	RPPEPL-LGAPQADASEEGVQRALDFAVSYNKGSDAYHRAIQWVRARQLVAGINY	69
Oy	68	FIDFVARETTCSKESNELTSC---ETKKLGOSLDCNAEYVYVPE 111	
Db	70	YLDVENGRTTCK-SQTNLT-NCPPHDQPHLRKALCSFOYKSVPEWK 114	

Search completed: September 24, 2004, 14:10:49
 Job time : 14.716 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: September 24, 2004, 14:08:41, Search time 44.704 Seconds
(without alignments)
913.519 Million cell updates/sec

Title: US-10-661-784-3

Perfect score: 687

Sequence: 1 GSGKDFVQPTKICVCPD.....VPMKKIYTVTVNNECEP 127

Scoring table: BLOSUM62

Gapop 10.0, Gapext 0.5

Searched: 1349238 seqs, 321558718 residues

Total number of hits satisfying chosen parameters: 1349238

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: Published Applications AA:

```
1: /cgn2_6/ptodata/1/pubpaas/US07_PUBCOMB.pep.*
2: /cgn2_6/ptodata/1/pubpaas/PCT_NEW_PUB.pep.*
3: /cgn2_6/ptodata/1/pubpaas/US06_NEW_PUB.pep.*
4: /cgn2_6/ptodata/1/pubpaas/US06_PUBCOMB.pep.*
5: /cgn2_6/ptodata/1/pubpaas/US07_NEW_PUB.pep.*
6: /cgn2_6/ptodata/1/pubpaas/PCTUS_PUBCOMB.pep.*
7: /cgn2_6/ptodata/1/pubpaas/US08_NEW_PUB.pep.*
8: /cgn2_6/ptodata/1/pubpaas/US08_PUBCOMB.pep.*
9: /cgn2_6/ptodata/1/pubpaas/US09A_PUBCOMB.pep.*
10: /cgn2_6/ptodata/1/pubpaas/US09B_PUBCOMB.pep.*
11: /cgn2_6/ptodata/1/pubpaas/US09C_PUBCOMB.pep.*
12: /cgn2_6/ptodata/1/pubpaas/US09_NEW_PUB.pep.*
13: /cgn2_6/ptodata/1/pubpaas/US10A_PUBCOMB.pep.*
14: /cgn2_6/ptodata/1/pubpaas/US10B_PUBCOMB.pep.*
15: /cgn2_6/ptodata/1/pubpaas/US10C_PUBCOMB.pep.*
16: /cgn2_6/ptodata/1/pubpaas/US10_NEW_PUB.pep.*
17: /cgn2_6/ptodata/1/pubpaas/US60_NEW_PUB.pep.*
18: /cgn2_6/ptodata/1/pubpaas/US60_PUBCOMB.pep.*
```

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	618	90.0	390	15	US-10-162-335-82
2	618	90.0	398	15	US-10-162-335-70
3	618	90.0	427	10	US-09-919-039-29
4	618	90.0	615	15	US-10-162-335-72
5	618	90.0	644	15	US-10-162-335-74
6	618	90.0	644	15	US-10-162-335-84
7	381	55.5	424	14	US-10-316-253-217
8	380	55.3	430	14	US-10-316-253-215
9	163.5	23.8	145	14	US-10-329-428-2
10	163.5	23.8	167	10	US-09-746-783-197
11	163	23.7	178	9	US-09-969-834-1
12	138.5	20.2	121	9	US-09-775-932-14
13	138.5	20.2	128	9	US-09-775-932-12
14	138.5	20.2	149	9	US-09-940-497-2
15	137.5	20.0	112	8	US-08-849-303-16

16	137.5	20.0	112	16	US-10-655-136-16
17	136.5	19.9	118	9	US-09-775-932-24
18	132.5	19.3	120	9	US-09-775-932-2
19	132.5	19.3	120	16	US-10-695-194-2
20	132.5	19.3	146	8	US-08-849-303-17
21	132.5	19.3	146	9	US-09-940-497-3
22	132.5	19.3	146	9	US-09-969-834-3
23	132.5	19.3	146	14	US-10-329-428-3
24	132.5	19.3	146	14	US-10-376-564-47
25	132.5	19.3	146	16	US-10-655-136-17
26	132.5	19.3	146	16	US-10-695-194-1
27	132.5	19.3	249	16	US-10-257-384A-4
28	132.5	19.3	641	16	US-10-257-384A-2
29	131.5	19.1	317	12	US-10-210-172-86
30	131.5	19.1	345	12	US-10-210-172-86
31	131.5	19.1	356	12	US-10-210-172-84
32	131.5	19.1	369	12	US-10-210-172-78
33	131.5	19.1	369	12	US-10-210-172-80
34	131.5	19.1	382	12	US-10-315-664-93
35	131.5	19.1	382	12	US-09-978-360A-425
36	130	18.9	127	8	US-08-849-303-19
37	130	18.9	127	16	US-10-655-136-19
38	129.5	18.9	140	14	US-10-376-564-46
39	129.5	18.9	140	14	US-10-376-564-48
40	128	18.6	111	8	US-08-849-303-26
41	128	18.6	111	16	US-10-655-136-26
42	127.5	18.6	140	8	US-08-849-303-18
43	127.5	18.6	140	16	US-10-655-136-18
44	124.5	18.1	121	9	US-09-775-932-8
45	124.5	18.1	141	8	US-08-849-303-24

ALIGNMENTS

RESULT 1

US-10-162-335-82

/ Sequence 82, Application US/10162335

/ Publication No. US20040009480A1

/ GENERAL INFORMATION:

/ APPLICANT: Anderson, David W.

/ APPLICANT: Baumgartner, Jason C.

/ APPLICANT: Boldog, Ferenc L.

/ APPLICANT: Casman, Stacie J.

/ APPLICANT: Edinger, Shlomit R.

/ APPLICANT: Gangolli, Esha A.

/ APPLICANT: Gerlach, Valerie

/ APPLICANT: Gorman, Linda

/ APPLICANT: Guo, Xiaojia (Sasha)

/ APPLICANT: Hsiao, Tord

/ APPLICANT: Kakuda, Ramesh

/ APPLICANT: Li, Li

/ APPLICANT: MacDougall, John R.

/ APPLICANT: Malyankar, Uriel M.

/ APPLICANT: Millet, Isabelle

/ APPLICANT: Padigaru, Muralidhara

/ APPLICANT: Patturajan, Meera

/ APPLICANT: Pena, Carol E. A.

/ APPLICANT: Rastelli, Luca

/ APPLICANT: Shinketa, Richard A.

/ APPLICANT: Stone, David J.

/ APPLICANT: Spytek, Kimberly A.

/ APPLICANT: Vernet, Corine A. M.

/ APPLICANT: Voser, Edward Z.

/ APPLICANT: Zernhusen, Bryan D.

/ TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Met

/ FILE REFERENCE: 21402-377 B

/ CURRENT APPLICATION NUMBER: US/10/162,335

/ PRIOR FILING DATE: 2002-10-01

/ PRIOR APPLICATION NUMBER: 60/295,607

/ PRIOR FILING DATE: 2001-06-04

/ PRIOR APPLICATION NUMBER: 60/295,651

/ PRIOR FILING DATE: 2001-06-04

```
/ PRIOR APPLICATION NUMBER: 60/296,404
/ PRIOR FILING DATE: 2001-06-06
/ PRIOR APPLICATION NUMBER: 60/296,418
/ PRIOR FILING DATE: 2001-06-06
/ PRIOR APPLICATION NUMBER: 60/297,414
/ PRIOR FILING DATE: 2001-06-11
/ PRIOR APPLICATION NUMBER: 60/297,567
/ PRIOR FILING DATE: 2001-06-12
/ PRIOR APPLICATION NUMBER: 60/298,285
/ PRIOR FILING DATE: 2001-06-14
/ PRIOR APPLICATION NUMBER: 60/298,556
/ PRIOR FILING DATE: 2001-06-15
/ PRIOR APPLICATION NUMBER: 60/299,949
/ PRIOR FILING DATE: 2001-06-21
/ PRIOR APPLICATION NUMBER: 60/300,883
/ PRIOR FILING DATE: 2001-06-26
/ Remaining Prior Application data removed - See File Wrapper or PALM.
/ NUMBER OF SEQ ID NOS: 201
/ SEQ ID NO 82
/ LENGTH: 390
/ TYPE: PRT
/ ORGANISM: Homo sapiens
US-10-162-335-82
```

```
Query Match          90.0%; Score 618; DB 15; Length 390;
Best Local Similarity 100.0%; Pred. No. 5,7e-59;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVCGPRDIPITNSPELEETLTHITKLAENNAATFYFKIDNVKARVQV 62
DB 216 GKDFVQPTKICVCGPRDIPITNSPELEETLTHITKLAENNAATFYFKIDNVKARVQV 275

QY 63 AGKKYFIDFVARETTCSKESNEELTSCETKLGOSLDCNAEVVVPWEKKIYPTV 118
DB 276 AGKKYFIDFVARETTCSKESNEELTSCETKLGOSLDCNAEVVVPWEKKIYPTV 331
```

```
RESULT 2
US-10-162-335-70
/ Sequence 70, Application US/10162335
/ Publication No. US2004009480A1
/ GENERAL INFORMATION:
/ APPLICANT: Anderson, David W.
/ APPLICANT: Baumgartner, Jason C.
/ APPLICANT: Boldog, Ferenc L.
/ APPLICANT: Casman, Stacie J.
/ APPLICANT: Edinger, Shlomit R.
/ APPLICANT: Gangoli, Zasha A.
/ APPLICANT: Gerlach, Valerie
/ APPLICANT: Gorman, Linda
/ APPLICANT: Guo, Xieojia (Sasha)
/ APPLICANT: Hjalt, Tord
/ APPLICANT: Kekuda, Ramesh
/ APPLICANT: Li, Li
/ APPLICANT: MacDougall, John R.
/ APPLICANT: Malyankar, Uriel M.
/ APPLICANT: Millet, Isabelle
/ APPLICANT: Padigaru, Muralidhara
/ APPLICANT: Parturajan, Meera
/ APPLICANT: Pena, Carol E. A.
/ APPLICANT: Restelli, Luca
/ APPLICANT: Shinketsu, Richard A.
/ APPLICANT: Stone, David J.
/ APPLICANT: Spytek, Kimberly A.
/ APPLICANT: Varnet, Corine A. M.
/ APPLICANT: Voss, Edward Z.
/ APPLICANT: Zernhusen, Bryan D.
/ TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
/ FILE REFERENCE: 21402-377 B
/ CURRENT APPLICATION NUMBER: US/10162,335
/ PRIOR FILING DATE: 2002-10-01
/ PRIOR APPLICATION NUMBER: 60/295,607
/ PRIOR FILING DATE: 2001-06-04
```

```
/ PRIOR APPLICATION NUMBER: 60/295,661
/ PRIOR FILING DATE: 2001-06-04
/ PRIOR APPLICATION NUMBER: 60/296,404
/ PRIOR FILING DATE: 2001-06-06
/ PRIOR APPLICATION NUMBER: 60/296,418
/ PRIOR FILING DATE: 2001-06-06
/ PRIOR APPLICATION NUMBER: 60/297,414
/ PRIOR FILING DATE: 2001-06-11
/ PRIOR APPLICATION NUMBER: 60/297,567
/ PRIOR FILING DATE: 2001-06-12
/ PRIOR APPLICATION NUMBER: 60/298,285
/ PRIOR FILING DATE: 2001-06-14
/ PRIOR APPLICATION NUMBER: 60/298,556
/ PRIOR FILING DATE: 2001-06-15
/ PRIOR APPLICATION NUMBER: 60/299,949
/ PRIOR FILING DATE: 2001-06-21
/ PRIOR APPLICATION NUMBER: 60/300,883
/ PRIOR FILING DATE: 2001-06-26
/ Remaining Prior Application data removed - See File Wrapper or PALM.
/ NUMBER OF SEQ ID NOS: 201
/ SEQ ID NO 70
/ LENGTH: 398
/ TYPE: PRT
/ ORGANISM: Homo sapiens
US-10-162-335-70

Query Match          90.0%; Score 618; DB 15; Length 398;
Best Local Similarity 100.0%; Pred. No. 5,8e-59;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVCGPRDIPITNSPELEETLTHITKLAENNAATFYFKIDNVKARVQV 62
DB 224 GKDFVQPTKICVCGPRDIPITNSPELEETLTHITKLAENNAATFYFKIDNVKARVQV 283

QY 63 AGKKYFIDFVARETTCSKESNEELTSCETKLGOSLDCNAEVVVPWEKKIYPTV 118
DB 284 AGKKYFIDFVARETTCSKESNEELTSCETKLGOSLDCNAEVVVPWEKKIYPTV 339
```

```
RESULT 3
US-09-919-039-29
/ Sequence 29, Application US/09919039
/ Publication No. US20030108871A1
/ GENERAL INFORMATION:
/ APPLICANT: Kasai, Matthew R.
/ TITLE OF INVENTION: GENES EXPRESSED IN TREATED HUMAN C3A LIVER CELL CULTURES
/ FILE REFERENCE: PA-0035 US
/ CURRENT APPLICATION NUMBER: US/09/919,039
/ PRIOR FILING DATE: 2002-09-09
/ PRIOR APPLICATION NUMBER: 60/222,113
/ PRIOR FILING DATE: 2000-07-28
/ NUMBER OF SEQ ID NOS: 401
/ SOFTWARE: PERL Program
/ SEQ ID NO 29
/ LENGTH: 427
/ TYPE: PRT
/ ORGANISM: Homo sapiens
/ FEATURE:
/ NAME/KEY: misc feature
/ OTHER INFORMATION: Incyte ID No. US20030108871A1 167507CD1
US-09-919-039-29
```

```
Query Match          90.0%; Score 618; DB 10; Length 427;
Best Local Similarity 100.0%; Pred. No. 6,4e-59;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 GKDFVQPTKICVCGPRDIPITNSPELEETLTHITKLAENNAATFYFKIDNVKARVQV 62
DB 253 GKDFVQPTKICVCGPRDIPITNSPELEETLTHITKLAENNAATFYFKIDNVKARVQV 312

QY 63 AGKKYFIDFVARETTCSKESNEELTSCETKLGOSLDCNAEVVVPWEKKIYPTV 118
DB 313 AGKKYFIDFVARETTCSKESNEELTSCETKLGOSLDCNAEVVVPWEKKIYPTV 368
```

DB 284 AGKYPIDFVARSTTCKESBELTSCETKKUGSLDCNAEVVWPKIPTV 339

RESULT 5
US-10-162-335-74
Sequence 74, Application US/10162335
Publication No. US20040009480A1
GENERAL INFORMATION:
APPLICANT: Anderson, David W.
APPLICANT: Baumgartner, Jason C.
APPLICANT: Boldog, Ferenc L.
APPLICANT: Casman, Stacie J.
APPLICANT: Edinger, Shlomit R.
APPLICANT: Gangoli, Esha A.
APPLICANT: Gerlach, Valerie
APPLICANT: Gorman, Linda
APPLICANT: Guo, Xiaojia (Sasha)
APPLICANT: Hjal, Tord
APPLICANT: Hjal, Tord
APPLICANT: Kekuda, Ramesh
APPLICANT: Li, Li
APPLICANT: MacDougall, John R.
APPLICANT: Malyankar, Uriel M.
APPLICANT: Millet, Isabelle
APPLICANT: Padigaru, Muralidhara
APPLICANT: Patturajan, Meera
APPLICANT: Pena, Carol E. A.
APPLICANT: Rastelli, Luca
APPLICANT: Shinkets, Richard A.
APPLICANT: Stone, David J.
APPLICANT: Spytek, Kimberly A.
APPLICANT: Vernet, Corine A. M.
APPLICANT: Voss, Edward Z.
APPLICANT: Zerhusen, Bryan D.
TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
FILE REFERENCE: 21402-377 B
CURRENT APPLICATION NUMBER: US/10/162,335
CURRENT FILING DATE: 2002-10-01
PRIOR APPLICATION NUMBER: 60/295,607
PRIOR FILING DATE: 2001-06-04
PRIOR APPLICATION NUMBER: 60/295,661
PRIOR FILING DATE: 2001-06-04
PRIOR APPLICATION NUMBER: 60/296,404
PRIOR FILING DATE: 2001-06-06
PRIOR APPLICATION NUMBER: 60/296,418
PRIOR FILING DATE: 2001-06-06
PRIOR APPLICATION NUMBER: 60/297,414
PRIOR FILING DATE: 2001-06-11
PRIOR APPLICATION NUMBER: 60/297,567
PRIOR FILING DATE: 2001-06-12
PRIOR APPLICATION NUMBER: 60/298,285
PRIOR FILING DATE: 2001-06-14
PRIOR APPLICATION NUMBER: 60/298,556
PRIOR FILING DATE: 2001-06-15
PRIOR APPLICATION NUMBER: 60/299,949
PRIOR FILING DATE: 2001-06-21
PRIOR APPLICATION NUMBER: 60/300,883
PRIOR FILING DATE: 2001-06-26
Remaining Prior Application data removed - See File Wrapper or PALM.
NUMBER OF SEQ ID NOS: 201
SEQ ID NO 74
LENGTH: 644
TYPE: PRT
ORGANISM: Homo sapiens
US-10-162-335-74

Query Match 90.0%; Score 618; DB 15; Length 644;
Best Local Similarity 100.0%; Pred. No. 1.le-58;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 GKDFVQPTKICVGCPRDPTNSPELETLTHITKLNANNATFYFKIDNVKARVQV 62
DB 253 GKDFVQPTKICVGCPRDPTNSPELETLTHITKLNANNATFYFKIDNVKARVQV 312

RESULT 4
US-10-162-335-72
Sequence 72, Application US/10162335
Publication No. US20040009480A1
GENERAL INFORMATION:
APPLICANT: Anderson, David W.
APPLICANT: Baumgartner, Jason C.
APPLICANT: Boldog, Ferenc L.
APPLICANT: Casman, Stacie J.
APPLICANT: Edinger, Shlomit R.
APPLICANT: Gangoli, Esha A.
APPLICANT: Gerlach, Valerie
APPLICANT: Gorman, Linda
APPLICANT: Guo, Xiaojia (Sasha)
APPLICANT: Hjal, Tord
APPLICANT: Hjal, Tord
APPLICANT: Kekuda, Ramesh
APPLICANT: Li, Li
APPLICANT: MacDougall, John R.
APPLICANT: Malyankar, Uriel M.
APPLICANT: Millet, Isabelle
APPLICANT: Padigaru, Muralidhara
APPLICANT: Patturajan, Meera
APPLICANT: Pena, Carol E. A.
APPLICANT: Rastelli, Luca
APPLICANT: Shinkets, Richard A.
APPLICANT: Stone, David J.
APPLICANT: Spytek, Kimberly A.
APPLICANT: Vernet, Corine A. M.
APPLICANT: Voss, Edward Z.
APPLICANT: Zerhusen, Bryan D.
TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
FILE REFERENCE: 21402-377 B
CURRENT APPLICATION NUMBER: US/10/162,335
CURRENT FILING DATE: 2002-10-01
PRIOR APPLICATION NUMBER: 60/295,607
PRIOR FILING DATE: 2001-06-04
PRIOR APPLICATION NUMBER: 60/295,661
PRIOR FILING DATE: 2001-06-04
PRIOR APPLICATION NUMBER: 60/296,404
PRIOR FILING DATE: 2001-06-06
PRIOR APPLICATION NUMBER: 60/296,418
PRIOR FILING DATE: 2001-06-06
PRIOR APPLICATION NUMBER: 60/297,414
PRIOR FILING DATE: 2001-06-11
PRIOR APPLICATION NUMBER: 60/297,567
PRIOR FILING DATE: 2001-06-12
PRIOR APPLICATION NUMBER: 60/298,285
PRIOR FILING DATE: 2001-06-14
PRIOR APPLICATION NUMBER: 60/298,556
PRIOR FILING DATE: 2001-06-15
PRIOR APPLICATION NUMBER: 60/299,949
PRIOR FILING DATE: 2001-06-21
PRIOR APPLICATION NUMBER: 60/300,883
PRIOR FILING DATE: 2001-06-26
Remaining Prior Application data removed - See File Wrapper or PALM.
NUMBER OF SEQ ID NOS: 201
SEQ ID NO 72
LENGTH: 615
TYPE: PRT
ORGANISM: Homo sapiens
US-10-162-335-72

Query Match 90.0%; Score 618; DB 15; Length 615;
Best Local Similarity 100.0%; Pred. No. 1e-58;
Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 GKDFVQPTKICVGCPRDPTNSPELETLTHITKLNANNATFYFKIDNVKARVQV 62
DB 254 GKDFVQPTKICVGCPRDPTNSPELETLTHITKLNANNATFYFKIDNVKARVQV 283
QY 63 AGKYPIDFVARSTTCKESBELTSCETKKUGSLDCNAEVVWPKIPTV 118

QY 63 AGKKYFIDFVARETTCSKESNEELTSCETKKGSLDCAAEVTVVWPKKIYPTV 118
 DB 313 AGKKYFIDFVARETTCSKESNEELTSCETKKGSLDCAAEVTVVWPKKIYPTV 368

RESULT 6

US-10-162-335-84
 ; Sequence 84, Application US/10162335
 ; Publication No. US2004000980A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Anderson, David W.
 ; APPLICANT: Baumgartner, Jason C.
 ; APPLICANT: Boldog, Ferenc L.
 ; APPLICANT: Casman, Stacie J.
 ; APPLICANT: Edinger, Shlomit R.
 ; APPLICANT: Gangolli, Esha A.
 ; APPLICANT: Gerlach, Valerie
 ; APPLICANT: German, Linda
 ; APPLICANT: Guo, Xiaojia (Sasha)
 ; APPLICANT: Hjalte, Tord
 ; APPLICANT: Kekuda, Rameesh
 ; APPLICANT: Li, Li
 ; APPLICANT: MacDougall, John R.
 ; APPLICANT: Malyankar, Uriel M.
 ; APPLICANT: Millet, Isabelle
 ; APPLICANT: Padigaru, Muralidhara
 ; APPLICANT: Patturajan, Meera
 ; APPLICANT: Pena, Carol E. A.
 ; APPLICANT: Rastelli, Luca
 ; APPLICANT: Shimkete, Richard A.
 ; APPLICANT: Stone, David J.
 ; APPLICANT: Spytek, Kimberly A.
 ; APPLICANT: Vernet, Corine A. M.
 ; APPLICANT: Voss, Edward Z.
 ; APPLICANT: Zetser, Bryan D.
 ; TITLE OF INVENTION: Therapeutic Polypeptides, Nucleic Acids Encoding Same, and Method
 ; FILE REFERENCE: 2402-377 B
 ; CURRENT APPLICATION NUMBER: US/10/162,335
 ; PRIOR FILING DATE: 2002-10-01
 ; PRIOR APPLICATION NUMBER: 60/295,607
 ; PRIOR FILING DATE: 2001-06-04
 ; PRIOR APPLICATION NUMBER: 60/295,661
 ; PRIOR FILING DATE: 2001-06-04
 ; PRIOR APPLICATION NUMBER: 60/296,404
 ; PRIOR FILING DATE: 2001-06-06
 ; PRIOR APPLICATION NUMBER: 60/296,418
 ; PRIOR FILING DATE: 2001-06-06
 ; PRIOR APPLICATION NUMBER: 60/297,414
 ; PRIOR FILING DATE: 2001-06-11
 ; PRIOR APPLICATION NUMBER: 60/297,567
 ; PRIOR FILING DATE: 2001-06-12
 ; PRIOR APPLICATION NUMBER: 60/298,285
 ; PRIOR FILING DATE: 2001-06-14
 ; PRIOR APPLICATION NUMBER: 60/298,556
 ; PRIOR FILING DATE: 2001-06-15
 ; PRIOR APPLICATION NUMBER: 60/299,949
 ; PRIOR FILING DATE: 2001-06-21
 ; PRIOR APPLICATION NUMBER: 60/300,883
 ; PRIOR FILING DATE: 2001-06-26
 ; Remaining Prior Application data removed - See File Wrapper or PALM.
 ; NUMBER OF SEQ ID NOS: 201
 ; SEQ ID NO 84
 ; LENGTH: 644
 ; TYPE: PRT
 ; ORGANISM: Homo sapiens
 US-10-162-335-84

Query Match 90.0%; Score 618; DB 15; Length 644;
 Best Local Similarity 100.0%; Pred. No. 1.1e-58;
 Matches 116; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVCGPRDPTNSPELEETLTHITTKLAENNAFFYFKIDNVKARQV 62

DB 253 GKDFVQPTKICVCGPRDPTNSPELEETLTHITTKLAENNAFFYFKIDNVKARQV 312
 QY 63 AGKKYFIDFVARETTCSKESNEELTSCETKKGSLDCAAEVTVVWPKKIYPTV 118
 DB 313 AGKKYFIDFVARETTCSKESNEELTSCETKKGSLDCAAEVTVVWPKKIYPTV 368

RESULT 7

US-10-316-253-217
 ; Sequence 217, Application US/10316253
 ; Publication No. US20030162706A1
 ; GENERAL INFORMATION:
 ; APPLICANT: The Procter & Gamble Company
 ; APPLICANT: Peters, Kevin
 ; APPLICANT: Thompson, Larry
 ; APPLICANT: Wang, Peng
 ; APPLICANT: Greis, Kenneth
 ; TITLE OF INVENTION: Angiogenesis Modulating Proteins
 ; FILE REFERENCE: 8865M
 ; CURRENT APPLICATION NUMBER: US/10/316,253
 ; CURRENT FILING DATE: 2002-12-10
 ; PRIOR APPLICATION NUMBER: US 60/355,295
 ; PRIOR FILING DATE: 2002-02-08
 ; NUMBER OF SEQ ID NOS: 308
 ; SOFTWARE: PatentIn version 3.1
 ; SEQ ID NO 217
 ; LENGTH: 424
 ; TYPE: PRT
 ; ORGANISM: Rattus norvegicus
 US-10-316-253-217

Query Match 55.5%; Score 381; DB 14; Length 424;
 Best Local Similarity 62.1%; Pred. No. 5.8e-33;
 Matches 72; Conservative 14; Mismatches 30; Indels 0; Gaps 0;
 QY 3 GKDFVQPTKICVCGPRDPTNSPELEETLTHITTKLAENNAFFYFKIDNVKARQV 62
 DB 246 GDDLPELLPNCRCGPRIEIPVDSPELKEALGHSLAQLNAQHNHIFYFKIDTVKATQV 305
 QY 63 AGKKYFIDFVARETTCSKESNEELTSCETKKGSLDCAAEVTVVWPKKIYPTV 118
 DB 306 AGVIVVIEFIARETNCQSKSTELTADCTEKLHGLSLCNAVYWRPWENKVTFTV 361

RESULT 8

US-10-316-253-215
 ; Sequence 215, Application US/10316253
 ; Publication No. US20030162706A1
 ; GENERAL INFORMATION:
 ; APPLICANT: The Procter & Gamble Company
 ; APPLICANT: Peters, Kevin
 ; APPLICANT: Thompson, Larry
 ; APPLICANT: Wang, Peng
 ; APPLICANT: Greis, Kenneth
 ; TITLE OF INVENTION: Angiogenesis Modulating Proteins
 ; FILE REFERENCE: 8865M
 ; CURRENT APPLICATION NUMBER: US/10/316,253
 ; CURRENT FILING DATE: 2002-12-10
 ; PRIOR APPLICATION NUMBER: US 60/355,295
 ; PRIOR FILING DATE: 2002-02-08
 ; NUMBER OF SEQ ID NOS: 308
 ; SOFTWARE: PatentIn version 3.1
 ; SEQ ID NO 215
 ; LENGTH: 430
 ; TYPE: PRT
 ; ORGANISM: Rattus norvegicus
 US-10-316-253-215

Query Match 55.3%; Score 380; DB 14; Length 430;
 Best Local Similarity 62.1%; Pred. No. 7.7e-33;
 Matches 72; Conservative 14; Mismatches 30; Indels 0; Gaps 0;


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QY 3 GDFVQPTKICVGCPRDIPNTSPBELTHTITKLNNAENNAFFKIDNVKARVQV 62
DB 252 GDFVQPTKICVGCPRDIPNTSPBELTHTITKLNNAENNAFFKIDNVKARVQV 311
QY 63 AGKXYPIDFVARETTCKSKESNEELTESECTKLGQSLDCNAEYVYVWPKKIYPTV 118
DB 312 AGVYVIEPIARETCKSKQSKTELTADECTKHLGQSLDCNANVYRPNKVKVPTV 367

RESULT 9
US-10-329-428-2
; Sequence 2, Application US/10329428
; Publication No. US20030114646A1
; GENERAL INFORMATION:
; APPLICANT: Li, et al.
; TITLE OF INVENTION: Human Cystatin P
; FILE REFERENCE: P2265PID2
; CURRENT APPLICATION NUMBER: US/10/329,428
; PRIOR FILING DATE: 2002-12-27
; PRIOR APPLICATION NUMBER: 60/014,795
; PRIOR FILING DATE: 1996-04-03
; PRIOR APPLICATION NUMBER: 08/832,535
; PRIOR FILING DATE: 1997-04-03
; PRIOR APPLICATION NUMBER: 09/019,485
; PRIOR FILING DATE: 1998-01-29
; PRIOR APPLICATION NUMBER: 09/528,436
; PRIOR FILING DATE: 2000-03-17
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: Patent in version 3.2
; SEQ ID NO 2
; LENGTH: 145
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-329-428-2

Query Match 23.8%; Score 163.5; DB 14; Length 145;
Best Local Similarity 31.6%; Pred. No. 9.7e-10;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

QY 11 TKICVGCPRDIPNTSPBELTHTITKLNNAENNAFFKIDNVKARVQVAGKXYPID 70
DB 32 SRVKGPPKTKINDPGVLOAARYSVKFNCTNDMLFKESRITRALVQIVKGLKYMLE 91
QY 71 FVARETTCKSKESNEELTESECTKLGQSLDCNAEYVYVWPKKIYPTVVAHWE 124
DB 92 VEIGRTTCKNOHLRL-DDCDFQTNHTLKQTLSCYSEVWVWVPW-----LQHFE 139

RESULT 10
US-09-746-783-197
; Sequence 197, Application US/09746783
; Publication No. US20030044935A1
; GENERAL INFORMATION:
; APPLICANT: Jacobs, Kenneth
; McCoy, John M.
; Lavalie, Edward R.
; Racie, Lisa A.
; Treacy, Maurice
; Spaulding, Vikki
; Agostino, Michael J.
; Howes, Steven H.
; Fectel, Kim
; TITLE OF INVENTION: SECRETED PROTEINS AND POLYNUCLEOTIDES
; ENCODING THEM
; NUMBER OF SEQUENCES: 231
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genetics Institute, Inc.
; STREET: 87 Cambridgepark Drive
; CITY: Cambridge
; STATE: MA
; COUNTRY: U.S.A.
; ZIP: 02140
; COMPUTER READABLE FORM:

QY 3 GDFVQPTKICVGCPRDIPNTSPBELTHTITKLNNAENNAFFKIDNVKARVQV 62
DB 252 GDFVQPTKICVGCPRDIPNTSPBELTHTITKLNNAENNAFFKIDNVKARVQV 311
QY 63 AGKXYPIDFVARETTCKSKESNEELTESECTKLGQSLDCNAEYVYVWPKKIYPTV 118
DB 312 AGVYVIEPIARETCKSKQSKTELTADECTKHLGQSLDCNANVYRPNKVKVPTV 367

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA: US/09/746,783
APPLICATION NUMBER: US/09/746,783
FILING DATE: 21-Dec-2000
CLASSIFICATION: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Milasincic, Debra J.
REGISTRATION NUMBER: 46,931
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 227-7400
TELEFAX: (617) 742-4214
INFORMATION FOR SEQ ID NO: 197:
SEQUENCE CHARACTERISTICS:
LENGTH: 167 amino acids
TYPE: amino acid
STRANDNESS: <Unknown>
TOPOLOGY: linear
MOLECULAR TYPE: Protein
SEQUENCE DESCRIPTION: SEQ ID NO: 197:
US-09-746-783-197

Query Match 23.8%; Score 163.5; DB 10; Length 167;
Best Local Similarity 31.6%; Pred. No. 1.2e-09;
Matches 37; Conservative 22; Mismatches 45; Indels 13; Gaps 3;

QY 11 TKICVGCPRDIPNTSPBELTHTITKLNNAENNAFFKIDNVKARVQVAGKXYPID 70
DB 54 SRVKGPPKTKINDPGVLOAARYSVKFNCTNDMLFKESRITRALVQIVKGLKYMLE 113
QY 71 FVARETTCKSKESNEELTESECTKLGQSLDCNAEYVYVWPKKIYPTVVAHWE 124
DB 114 VEIGRTTCKNOHLRL-DDCDFQTNHTLKQTLSCYSEVWVWVPW-----LQHFE 160

RESULT 11
US-09-969-834-1
; Sequence 1, Application US/09969834
; Patent No. US20020102711A1
; GENERAL INFORMATION:
; APPLICANT: Bandman, Olga
; Golli, Surya K.
; TITLE OF INVENTION: NOVEL HUMAN CYSTATIN-LIKE
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/969,834
; FILING DATE: 01-Oct-2001
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/09/471,765
; FILING DATE: <Unknown>
; APPLICATION NUMBER: US/08/791,522
; FILING DATE: <Unknown>
; APPLICATION NUMBER: 09/471,765
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
```

; REFERENCE/DOCKET NUMBER: PP-0193 US
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: 415-895-0555
 ; FAX: 415-845-4166
 ; INFORMATION FOR SEQ ID NO: 1:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 178 amino acids
 ; TYPE: amino acid
 ; STRANDEDNESS: single
 ; TOPOLOGY: linear
 ; IMMEDIATE SOURCE:
 ; CLONE: 30443
 ; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
 US-09-969-834-1

Query Match 23.78; Score 163; DB 9; Length 178;
 Best Local Similarity 34.04; Pred. No. 1.4e-09;
 Matches 35; Conservative 20; Mismatches 44; Indels 4; Gaps 2;
 QY 11 TKICVGCPRDIPNTSPLEETLTHITIKLAENNAATFYFKIDNVKARVQVAGKKYFI 70
 DB 54 SRVKGPFKTKNDPQVQAARVSYVEKFNCTNDMFLEKESRITRALVQIVKGLKYLE 113
 QY 71 FVARETTCSKE---SNEELTESCETKGLQ--SLDCNAEVVVPWE 110
 DB 114 VEIGRTCKGNQHLRL--DCDFQTNHTLKQTLSCYSEWVVPW 155

RESULT 12
 US-09-775-932-14
 ; Sequence 14, Application US/09775932
 ; Patent No. US20020137671A1
 ; GENERAL INFORMATION:
 ; APPLICANT: University of British Columbia
 ; TITLE OF INVENTION: Production and use of Modified Cystatins
 ; FILE REFERENCE: 58069
 ; CURRENT APPLICATION NUMBER: US/09/775,932
 ; CURRENT FILING DATE: 2001-02-02
 ; PRIOR APPLICATION NUMBER: CA99/00717
 ; PRIOR FILING DATE: 1999-08-05
 ; PRIOR APPLICATION NUMBER: 60/095,503
 ; PRIOR FILING DATE: 1998-08-05
 ; NUMBER OF SEQ ID NOS: 32
 ; SOFTWARE: Patentin Ver. 2.0
 ; SEQ ID NO 14
 ; LENGTH: 121
 ; TYPE: PRT
 ; ORGANISM: Homo sapiens
 US-09-775-932-14

Query Match 20.24; Score 138.5; DB 9; Length 121;
 Best Local Similarity 31.54; Pred. No. 4.2e-07;
 Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;
 QY 10 PTKICVGCPRDIPNTSPLEETLTHITIKLAENNAATFYFKIDNVKARVQVAGKKYFI 69
 DB 2 PQRMVWGLRDLSPDPQVQAARVSYVFNMGNSIYFRDTHIIKAQSQLVAGIKYFL 61
 QY 70 DFVARETTCSKE---SNEELTESCETKGLQ--SLDCNAEVVVPWE 111
 DB 62 TMWNGSTDCRKRTRVTDGHDVLT-TCPLAAGAQOEKLRCDFEVLVVPWQ 108

RESULT 13
 US-09-775-932-12
 ; Sequence 12, Application US/09775932
 ; Patent No. US20020137671A1
 ; GENERAL INFORMATION:
 ; APPLICANT: University of British Columbia
 ; TITLE OF INVENTION: Production and use of Modified Cystatins
 ; FILE REFERENCE: 58069
 ; CURRENT APPLICATION NUMBER: US/09/775,932
 ; CURRENT FILING DATE: 2001-02-02

; PRIOR APPLICATION NUMBER: CA99/00717
 ; PRIOR FILING DATE: 1999-08-05
 ; PRIOR APPLICATION NUMBER: 60/095,503
 ; PRIOR FILING DATE: 1998-08-05
 ; NUMBER OF SEQ ID NOS: 32
 ; SOFTWARE: Patentin Ver. 2.0
 ; SEQ ID NO 12
 ; LENGTH: 128
 ; TYPE: PRT
 ; ORGANISM: Homo sapiens
 US-09-775-932-12

Query Match 20.24; Score 138.5; DB 9; Length 128;
 Best Local Similarity 31.54; Pred. No. 4.5e-07;
 Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;
 QY 10 PTKICVGCPRDIPNTSPLEETLTHITIKLAENNAATFYFKIDNVKARVQVAGKKYFI 69
 DB 9 PQRMVWGLRDLSPDPQVQAARVSYVFNMGNSIYFRDTHIIKAQSQLVAGIKYFL 68
 QY 70 DFVARETTCSKE---SNEELTESCETKGLQ--SLDCNAEVVVPWE 111
 DB 69 TMWNGSTDCRKRTRVTDGHDVLT-TCPLAAGAQOEKLRCDFEVLVVPWQ 115

RESULT 14
 US-09-940-497-2
 ; Sequence 2, Application US/09940497
 ; Patent No. US20020052476A1
 ; GENERAL INFORMATION:
 ; APPLICANT: NI et al.
 ; TITLE OF INVENTION: Human Cystatin E
 ; FILE REFERENCE: PF202PID2
 ; CURRENT APPLICATION NUMBER: US/09/940,497
 ; CURRENT FILING DATE: 2001-08-29
 ; PRIOR APPLICATION NUMBER: US 09/241,376
 ; PRIOR FILING DATE: 1999-02-02
 ; PRIOR APPLICATION NUMBER: US 08/744,138
 ; PRIOR FILING DATE: 1996-11-05
 ; PRIOR APPLICATION NUMBER: US 08/461,030
 ; PRIOR FILING DATE: 1995-06-05
 ; NUMBER OF SEQ ID NOS: 13
 ; SOFTWARE: Patentin version 3.1
 ; SEQ ID NO 2
 ; LENGTH: 149
 ; TYPE: PRT
 ; ORGANISM: Homo sapiens
 US-09-940-497-2

Query Match 20.24; Score 138.5; DB 9; Length 149;
 Best Local Similarity 31.54; Pred. No. 5.5e-07;
 Matches 34; Conservative 22; Mismatches 45; Indels 7; Gaps 3;
 QY 10 PTKICVGCPRDIPNTSPLEETLTHITIKLAENNAATFYFKIDNVKARVQVAGKKYFI 69
 DB 30 PQRMVWGLRDLSPDPQVQAARVSYVFNMGNSIYFRDTHIIKAQSQLVAGIKYFL 89
 QY 70 DFVARETTCSKE---SNEELTESCETKGLQ--SLDCNAEVVVPWE 111
 DB 90 TMWNGSTDCRKRTRVTDGHDVLT-TCPLAAGAQOEKLRCDFEVLVVPWQ 136

RESULT 15
 US-08-849-303-16
 ; Sequence 16, Application US/08849303
 ; Publication No. US20030221209A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Atkinson, Howard J.
 ; APPLICANT: McPherson, Michael J.
 ; APPLICANT: Urwin, Peter E.
 ; TITLE OF INVENTION: MODIFIED PROTEINASE INHIBITORS
 ; NUMBER OF SEQUENCES: 79
 ; CORRESPONDENCE ADDRESS:

UB-10-661-784-3.rapb

Mon Sep 27 08:32:57 2004

ADDRESSEE: Klauber & Jackson
STREET: 411 Hackensack Avenue, 4th Floor
CITY: Hackensack
STATE: New Jersey
COUNTRY: USA
ZIP: 07601
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA: 303
APPLICATION NUMBER: US/08/849,303
FILING DATE: 21-MAY-1997
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Jackson Esq., David A.
REGISTRATION NUMBER: 26,742
REFERENCE/DOCKET NUMBER: 1321-1-003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 201-487-5800
TELEFAX: 201-343-1684
TELEX: 133521
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 112 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
HYPOTHETICAL: NO
US-08-849-303-16

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Query Match      20.0%; Score 137.5; DB 8; Length 112;
Best Local Similarity 28.8%; Pred.No 4.9e-07;
Matches 32; Conservative 25; Mismatch 35; Indels 19; Gaps 4;

24 NSPELBETLTHITKLNABNNATFYKIDNVKARQVQVAGKCYFDIFVARETTSKSGSN 83
      : : : : : : : : : : : : : : : : : : : : : : : : : : : :
12 NEEGVOEALSPAVSEPNRSGNDAYQERVVVRVRAKQVVSQMYFLDELGLRGTCTK--S 69
      : : : : : : : : : : : : : : : : : : : : : : : : : : : :

84 BELTESC-----ETKKLGSLDCNAEYVVPVPEKKIYFTVTNNHCEC 126
      : : : : : : : : : : : : : : : : : : : : : : : : : : : :

70 QAKLDCSPHNPQHLKREKL-----CSFQVYVVPWN-----TINLVFSCQ 111
      : : : : : : : : : : : : : : : : : : : : : : : : : : : :

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Job time : 45.704 secs

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